

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: July 19, 2002, 21:31:06 ; Search time 2528.22 Seconds
(without alignments)

18247.064 Million cell updates/sec

Title: US-09-836-410A-2
Perfect score: 3418
Sequence: 1 caagtaacacccgcaagatg.....atgcaataaattgttttggg 3418

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 13736207 seqs, 6748477542 residues

Total number of hits satisfying chosen parameters: 27472414

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:*

1: em_estba:*
2: em_esthum:*
3: em_estin:*
4: em_estnu:*
5: em_estov:*
6: em_estpl:*
7: em_estro:*
8: em_hic:*
9: gb_est1:*
10: gb_est2:*
11: gb_hic:*
12: gb_gss:*
13: em_gss_hum:*
14: em_gss_inv:*
15: em_gss_pln:*
16: em_gss_vrt:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	950.2	27.8	2758	11	AK005056 Mus muscu
2	754.4	22.1	1093	10	BM456794 AGENCOURT
3	714.2	20.9	1959	11	AK007755 Mus muscu
4	704.8	20.6	821	10	BM044197
5	679.2	19.9	704	10	BM067031
6	673.4	19.7	689	9	BB616617
7	656.4	19.2	659	9	BB659255
8	649.8	19.0	675	9	AV270853
9	646.2	18.9	652	9	AW107262
10	644.6	18.9	654	10	BF012472
11	629.6	18.4	673	9	BB577716
12	623.8	18.3	635	9	AW260482
13	623.4	18.2	625	10	BM080108
14	617.8	18.1	710	9	AI744486
15	616.2	18.0	719	9	AU130763
16	610	17.8	710	10	BF163411
17	608	17.8	617	9	BB478039

18	607.4	17.8	744	10	BG671648
19	602.2	17.6	945	10	BF179047
20	601	17.6	978	10	BM463359
21	591	17.3	657	10	BF472586
22	588.2	17.2	593	10	BG065941
23	580	17.0	600	10	BG805362
24	579	16.9	941	10	BG823888
25	575	16.8	603	10	BG079208
26	574.8	16.8	589	9	AW048763
27	573.4	16.8	626	10	BI662491
28	571.6	16.7	578	10	BI078250
29	561.4	16.4	629	10	BE300741
30	559.2	16.4	623	9	AW534169
31	559	16.4	625	10	BM236189
32	545.8	16.0	549	10	BE852629
33	545.2	16.0	684	9	AI177404
34	541.2	15.8	582	10	BM022195
35	539.6	15.8	615	10	BI966451
36	537	15.7	549	10	BF453530
37	535.2	15.7	931	10	BI687745
38	529.8	15.5	601	10	BF472327
39	528.8	15.5	666	10	D86662
40	518.8	15.2	572	11	AK017653
41	516.8	15.1	600	10	BI989826
42	507.8	14.9	519	10	BF450556
43	506.8	14.8	510	9	AA435048
44	505.4	14.8	826	10	BI150077
45	505.2	14.8	511	10	BE626970

ALIGNMENTS

AK005056 2758 bp mRNA linear HTC 19-JAN-2002
Mus musculus adult male liver cDNA, RIKEN full-length enriched library, clone:1300019C06:related to PUTATIVE N-TERMINAL ACETYLTRANSFERASE, full insert sequence.

AK005056
HTC; CAP trapper.
Mus musculus (strain:C57BL/6J) adult male liver cDNA to mRNA, clone:1300019C06.
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (sites)
Carninci, P. and Hayashizaki, Y.
High-efficiency full-length cDNA cloning
Meth. Enzymol. 303, 19-44 (1999)

REFERENCE 2 (sites)
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes
Genome Res. 10 (10), 1617-1630 (2000)

REFERENCE 3 (sites)
Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P., Konno, H., Akiyama, J., Nishi, K., Kitsumai, T., Tashiro, H., Itoh, M., Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Harada, A., Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K., Fujiwaki, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer
Genome Res. 10 (11), 1757-1771 (2000)

QY 887 tgaagccagccctggacacagcagacagatttattaataatccaagtgtgcaaaatacat 946
DB 1359 TGAGGCACAGTCTTTGGACACCGCTTGACAGGTTCTCAATTCCTCAATATGTCCTGAGGCTCTCTGATGGCAG 2498
QY 947 gtaaaagccaacctgattaaagagcgtgaagaaatgtgttccaagtttccaggggaagg 1006
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QY 1127 gagacattttatagaataaccagtagaccagtttgactttacatacatctgtatgagaa 1186
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QY 2087 ttatctgttggctttcgttcgtcctcctcctggatacagaag 2121
DB 2559 ACTCACACCTGCTTTCTCTGCTGCTGCTGCGAGGAAG 2593
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LOCUS AGENCOURT_6404070 NIH_MGC_92 Homo sapiens cDNA clone IMAGE:5583707
DEFINITION 5', mRNA sequence.
ACCESSION BM456794
VERSION BM456794.1 GI:18505834
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 1093)
AUTHORS NIH-MGC <http://mgc.nci.nih.gov/>.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
Plate: LLAM12347 row: d column: 12
High quality sequence stop: 695.
FEATURES
source
1. 1093
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:5583707"
/clone_1lb="NIH_MGC_92"
/tissue_type="embryonal carcinoma, cell line"
/lab_host="DH10B (phage-resistant)"
/note="Organ: testis; Vector: pCMV-SPORT6; Site_1: NotI;
Site_2: SalI; Cloned unidirectionally; oligo-dT primed.
Average insert size 2.5 kb. Library enriched for
full-length clones and constructed by Life Technologies.
Note: this is a NIH_MGC Library."
BASE COUNT 415 a 183 c 234 g 253 t 8 others
ORIGIN
Query Match 22.1%; Score 754.4; DB 10; Length 1093;
Best Local Similarity 91.5%; Pred. No. 5e-119;
Matches 820; Conservative 0; Mismatches 73; Indels 3; Gaps 2;
QY 683 tggaaagaggaaacctccaaccacattactttgggtccagctactatttggcacagcattta 742
DB 1 TGGAAAGAGGAAACCAACCAACCATTTACTTTGGTCCAGTACTTGGCACAACATTA 60
QY 743 tgataaattggtcagccatccattgtctctggaatacataataactgcaattgaaaglac 802
DB 61 TGACAAAATTTGGTCAGCCATCTATTGCTTTGGAGTACATAAATACTGCTATTGAAGTAC 120
QY 803 accaacattgatagaactctttctgttaaaagctaaactatataagcatgctgggaatat 862
DB 121 ACCTACATTAATGAAGCTCTTTCTCTGGAAGCTAAATCTATTAAGCATGCTGGAAATAT 180
QY 863 taaagaagctgccagggtggatggatgaagccaggcccttggaacagcagcagcagatttat 922

BASE COUNT 633 a 377 c 454 g 495 t
ORIGIN related to PUTATIVE N-TERMINAL ACETYLTRANSFERASE"

Query Match 20.9%; Score 714.2; DB 11; Length 1959;
Best Local Similarity 69.0%; Pred. No. 3e-112;
Matches 977; Conservative 0; Mismatches 438; Indels 0; Gaps 0;

QY 47 tggattgggtatgctttaccattattagaagactatgaaatggcagcaaaaattttaga 106
DB 545 TGGATATGCCATCGCTTATTCATTTGCTGAAGATTATGATACAGCGCTAAAGCTATTAGA 604

QY 107 agagtttagaaacacacagcagacatctctgataaagtggattatgaatagtaact 166
DB 605 AGAATTTAGACAACGCAACAGTTCCTCCCAACAAAATAGCTTTATGAATATAGTGAAT 664

QY 167 cctcttatcatcagaatcaagttcttcggaagcaggtctttatagagaagccttggaaca 226
DB 665 GTTACTATATCAGATCAAGTCATGAGAGAGGCAAAATTTATTTTCAGGAATCTTTGGAACA 724

QY 227 tctttgtacctatgaagcagatttgatataacttgcgttgtaagaacaccaaaggga 286
DB 725 TATGAACAACATAGAGAAGTTGATATGTGATAAAGCTTTTGGTGGGAAGAAATTAAGGGGA 784

QY 287 acttctgttcagttgctgctttggaagatgctgtaacgttttatagagattacaaga 346
DB 785 AATGCTGTTGAATTTGGGAGACTAAAGAACCCAGTGAAGTATTCAGAAACTTGATTGA 844

QY 347 gaggaatcctgaaattgggctattacaaggcttagaagaagcactgaagccagctaa 406
DB 845 TTGGAATGCTGAAAATTTGGTGTATTATATGAAGGCTTTGGAAGGCAATTCACACTTCGTTTC 904

QY 407 tatgttaaaagcgttaaaatatatagaagaagcctggactaaataccccaggggactcgt 466
DB 905 TCTAGACGAGGCTTCAGCTTTATGAGAAGGCTCAGCAGCAGCAGCAGCAGCAGCTTC 964

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QY 527 gtccctaaggatgaatttcagcaagggtgtccactgtcttcaataccttgaggtcttt 586
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QY 647 tctaaaaagttgtcgcctatttaaccccaatgatgaaaggaggaacccctcccaacc 706
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QY 1187 gatccctctgatcatcatgtggtgacttattaaaaacagagaatgtacttcagcagatcc 1246
DB 1685 GATGACCCCTCGCTGTATGTGTGGCTCTTGAGATTAGAAATGCTCTCAGAAGACATAC 1744

QY 1247 atttactctcaaacagcagagaaattgctattgagacttattgaagcttcatgacaaccc 1306
DB 1745 TTTTATTTCAGGCTGTAGATCAGCAATTTGAAATATATTTGAAATTTACATGATAACCC 1804

QY 1307 tctgacagatgagaacaaagacagagctgatacagacaaacatgtctgacaaagagct 1366
DB 1805 TTAAACCAATGACAGCAAAACAAACAGACATAGATTTCAGAAAAACCTGTGAGCAAGAAAT 1864

QY 1367 aaagaaactgcgttaataacaaagaagagctcaaaagaagcccgagattgaagaagaga 1426
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DB 1925 AAAGCACACAGAAAGCGAACGCCAACACAAAAAACC 1959

RESULT 4

BM044197 821 bp mRNA linear EST 07-NOV-2001
LOCUS 603621442F1 NIH_MGC_40 Homo sapiens CDNA clone IMAGE:5446979 5',
DEFINITION mRNA sequence.
ACCESSION BM044197
VERSION BM044197.1 GI:16773464
KEYWORDS EST.
SOURCE human.

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 821)
AUTHORS NIH-MGC http://mgc.nci.nih.gov/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgabbs-remail.nih.gov
Tissue Procurement: DCTD/DTP
CDNA Library Preparation: Ling Hong/Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LICM1930 row: c column: 12
High quality sequence stop: 820.

FEATURES
Source

1. 821
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/clone_image="IMAGE:5446979"
/clone_lib="NIH_MGC_40"
/tissue_type="carcinoma, cell line"
/lab_host="DH10B (phage-resistant)"
/note="Organ: prostate; Vector: pOTB7; Site_1: XhoI;
Site_2: EcoRI; cDNA made by oligo-dt priming.
Directionally cloned into EcoRI/XhoI sites using the
following 5' adaptor: GGCACAG(G). Library constructed by
Ling Hong in the laboratory of Gerald M. Rubin (University

of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies).
Note: this is a NIH_MGC Library.
BASE COUNT 310 a 145 c 168 g 198 t
ORIGIN

Query Match 20.6%; Score 704.8; DB 10; Length 821;
Best Local Similarity 92.8%; Pred. No. 1.5e-110; Indels 2; Gaps 2;
Matches 761; Conservative 0; Mismatches 57; Indels 2; Gaps 2;
QY 594 gataaagaagagtggtgcaactctgtagaagaacttagtggtgtatgaaacttctctaaaa 653
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QY 954 gcaaacctgtattaagaaggtggaagaattggttccaaagtattcagggggaagaactca 1013
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Db 782 CTTAGTATATAAACAAGAAAGAAAGCTCAAAAGAAAGGCCCA 821

RESULT 5
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LOCUS
DEFINITION H3049G08-3 NTA Mouse 15K cDNA Clone Set Mus musculus linear EST 26-JAN-2001
H3049G08 3', mRNA sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

REFERENCE,
AUTHORS
TITLE
JOURNAL
COMMENT

FEATURES
source

BG067031
BG067031.1 GI:12549600
EST.
house mouse.
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 704)
Kargul,G.J., Dudekula,D.B., Qian,Y., Lim,M.K., Jaradat,S.A., Tanaka
,T.S., Carter,M.G. and Ko,M.S.H.
Verification and initial annotation of NIA mouse 15K cDNA clone set
Unpublished (2001)
Other_ESTS: H3049G08-5
Contact: George J. Kargul
Laboratory of Genetics
National Institute on Aging/National Institutes of Health
333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6820, USA
Email: cdna@lgsun.grc.nia.nih.gov
This clone set has been freely distributed to the community. Please
visit <http://lgsun.grc.nia.nih.gov/cDNA/15k.html> for details.
Plate: H3049 row: G column: 08
Seq primer: -21M13 Forward
High quality sequence stop: 704
POLYA=tes.

Location/Qualifiers

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/strain="C57BL/6J"
/db_xref="niaEST:H3049G08-3"
/db_xref="taxon:10090"
/clone="H3049G08"
/clone_lib="NIA Mouse 15K cDNA Clone Set"
/sex="Clones arrayed from a variety of cDNA libraries"
/dev_stage="Clones arrayed from a variety of cDNA libraries"
/lab_host="DH10B"
/note="Vector: pSPORT1; Site_1: Salt; Site_2: NotI; This clone is among a rearranged set of 15,247 clones from 11 embryo cDNA libraries (including preimplantation stage embryos from unfertilized egg to blastocyst, embryonic part of E7.5 embryos, extraembryonic part of E7.5 embryos, and E12.5 female mesonephros/gonad) and one newborn ovary cDNA library. Average insert size 1.5 kb. All source libraries are cloned unidirectionally with Oligo(dT)-Not primers. References include: (1) Genome-wide expression profiling of mid-gestation placenta and embryo using a 15,000 mouse developmental cDNA microarray, 2000, Proc. Natl. Acad. Sci. U S A, 97: 9127-9132; (2) Large-scale cDNA analysis reveals phased gene expression patterns during preimplantation mouse development, 2000, Development, 127: 1737-1749; (3) Genome-wide mapping of unselected transcripts from extraembryonic tissue of 7.5-day mouse embryos reveals enrichment in the t-complex and under-representation on the X chromosome, 1998, Hum Mol Genet 7: 1967-1978."
Mol Genet 7: 1967-1978."

BASE COUNT 168 a 149 c 124 g 263 t

ORIGIN

Query Match 19.9%; Score 679.2; DB 10; Length 704;
Best Local Similarity 99.4%; Pred. No. 3.7e-106;
Matches 692; Conservative 0; Mismatches 3; Indels 1; Gaps 1;
QY 729 ttggcacagcattatgataaaattggtcagccattccattgtctctggaatacataact 788
Db 695 TTTGCACAGCATTTATGATAAAATTTGGTCAGCCATCCATTGCTCTGGAATACATAAACT 636
QY 789 gcaattgaaagtacacacacattgatagaactcttctgtgaaagctataaataaag 848
Db 635 GCAATTGAAAGTACACCAACATGATGAACTCTTTCTGTAAGAGCTAATCTAAG 576
QY 849 catgctgggaataataaagaagctgccagggtggatgatgaagcccgccctggacaca 908
|||||


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Db 421 ACCAACAGAAACCTTCAGACTTCGATGGAGGTGTTGGAAGCCCTTGTGTGATGGTAGCCTA 480
QY 2031 cgagactgtaaagaagctgcgaagcctacagagcaagtgctgtcataagcttttcccttat 2090
Db 481 GGAGAGCTGTAAAGAAGCTGCCGAAGCCTACAGAGCAAGTTGTGTAAGCTTTTCCCTTAT 540
QY 2091 gcttggtcttcacgctcctcctgatacagagagagatgaagatcacagtgcaacgagat 2150
Db 541 GCTTTGGCTTTTCATGCTCCTCGATACGAGAGGATATGAAGATCACAGTGCAACGGAGAT 600
QY 2151 agtctgcagaacggaagaaactgcccgaatgaaatctgacatcattaaacaagcaaatg 2210
Db 601 AGTTCTGAGAACGGAAGAACTGGCCATGAATCTGAACATCATTAACAAGCAAAATG 660
QY 2211 gaatgactttggacc 2225
Db 661 GAATGACTTTGGACC 675

RESULT 9
AW107262/c
LOCUS
DEFINITION
unl3c03.x1 Sugano mouse kidney mkia Mus musculus cDNA clone
IMAGE:2192164 3' similar to WP:Y50D7_164.A CE22298 ;, mRNA
sequence.
ACCESSION
AW107262
VERSION
AW107262.1 GI:6078062
KEYWORDS
EST.
SOURCE
house mouse.
ORGANISM
Mus musculus
REFERENCE
1 (bases 1 to 652)
AUTHORS
Marra,M., Hillier,L., Kucaba,T., Martin,J., Beck,C., Wylie,T.,
Underwood,K., Steptoe,M., Theising,B., Allen,M., Bowers,Y., Person
B., Swaller,T., Gibbons,M., Pape,D., Harvey,N., Schurk,R., Ritter
E., Kohn,S., Shin,T., Jackson,Y., Cardenas,M., McCann,R.,
Waterston,R. and Wilson,R.
The WashU-NCI Mouse EST Project 1999
Unpublished (1999)
TITLE
Marra M/WashU-NCI Mouse EST Project 1999
JOURNAL
Washington University School of Medicine
COMMENT
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1810
Email: mouseest@wustl.edu
This clone is available royalty-free through LBNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
MGI:1004616
Seq primer: custom primer used
High quality sequence stop: 493.
Location/Qualifiers
FEATURES
source
1..652
/organism="Mus musculus"
/strain="C57BL"
/db_xref="taxon:10090"
/clone="IMAGE:2192164"
/clone_lib="Sugano mouse kidney mkia"
/sex="female"
/dev_stage="adult"
/lab_host="DH10B"
/notes="Organ: kidney; Vector: pME18S-FL3; Site_1: DraIII
(CACTGTGTG); Site_2: DraIII (CAGCATGTG); 1st strand cDNA
was primed with an oligo(dT) primer
[ATGTGGCTTTTCTTTTCTTTT]; double-stranded cDNA was
ligated to a DraIII adaptor [TGTGGCTACTGG], digested
(CACTGTGTG); Site_2: DraIII (CAGCATGTG) sites of the pME18S-FL3
vector (5' site CACTGTGTG, 3' site CAGCATGTG). XhoI should
be used to isolate the cDNA insert. Size selection was
performed to exclude fragments <1.5kb. Library
constructed by Dr. Sumio Sugano (University of Tokyo
Institute of Medical Science). Custom primers for
sequencing: 5' end primer CTCTGTCTCTAAAGAGCTGCG and 3' end

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BASE COUNT 155 a 138 c 114 g 244 t 1 others
ORIGIN
Query Match 18.9%; Score 646.2; DB 9; Length 652;
Best Local Similarity 99.4%; Pred. No. 1.6e-100;
Matches 648; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
QY 765 attgctctggaatatacaataactcaattgaatcaccacacattgatgaactcttt 824
Db 652 ATTGCTCTGGAATACATAAATACTGCAATTGAAAGTACACCAACATTGATAGAACTCTT 593
QY 825 ctgttaaaagcttaaaatctataagcatgctgggaattattaaagaagctgcagggtggatg 884
Db 592 CTGTGTAAGAAGCTANAATCTATAAGCATGCTGGCAATATTAAAGAAGCTGCCAGGTGGATG 533
QY 885 gatgaagcccgccctggacacagacagacagatttattattccaaagtgtcaaaatc 944
Db 532 GATGAAGCCCGCCCTGGACACAGACAGACAGATTATTAATTCGAAGTGTGCAAAATAC 473
QY 945 atgttaaaagcccaactgattaaagagggctgaagaaatgtgttccaagtttacgagggaa 1004
Db 472 ATGTTAAAAGCCCACTGATTAAGAGAGGCTGAAGAAATGTGTTCGAAGTTTACGAGGGAA 413
QY 1005 ggaacttcagcggtagagaaacctgaatgaaatgcagtgatgtgtgttcagacagagtg 1064
Db 412 GGAACCTTCAGCGGTAGAGAACCTGAATGAATGCCAGTGTATGTGTTCAGACAGAGTGT 353
QY 1065 gctcaggcatacaagcaatgaacaaatgtgtgaagcacttaagaataatgtcatgaaatt 1124
Db 352 GCTCAGGCATACAAAGCAATGAACAAATTTGGTGAAGCACTTAAGAAATGTGATGAAT 293
QY 1125 gagagacatttatagaatacccgatgaccagtttgacccagtttgactttcactacatctgtatgag 1184
Db 292 GAGAGACATTTATAGAAATCACCAGTACCAGTGTGACTTTTCATACATCTGATGAGG 233
QY 1185 aagatcaccccttagatcatatgtggacttatataaactagaagatctacttcagacagat 1244
Db 232 AGATCACCCCTTAGATCATATGTGGACTTATTAAAACTAGAAGATGTACTTCGACAGCAT 173
QY 1245 ccattttactcaaacagcagcaaatgtctattgagatctatttgaagcttcatgacaac 1304
Db 172 CCATTTTACTTCAAGCAGCGAGGATTGCTATTGAGATCTATTGAGCTTTCATGACAAC 113
QY 1305 cctctgacagatgagacaagaacacagcaggctgtatcacgcaaacatgtctgacaagag 1364
Db 112 CCTCTGACAAATGAGAACAAAGAACACGAGGCTGATACAGCAACATGTCTGCACAAAGAG 53
QY 1365 ctaagaagaactgcgttaataaacaagaagagctcaagaagacccagattg 1416
Db 52 CTAAGAAGAACTGCGTAAATAAACAAGAAGAGCTCAAAAGAAAGCCCGCATAG 1

RESULT 10
BF012472
LOCUS
DEFINITION
ux56g03.y1 Soares_NKWMND_mandible Mus musculus cDNA clone
IMAGE:3514324 5' similar to TR:Q9VWI2_Q9VWI2 CGI2202 PROTEIN. ;,
mRNA sequence.
ACCESSION
BF012472
VERSION
BF012472.1 GI:10712747
KEYWORDS
EST.
SOURCE
house mouse.
ORGANISM
Mus musculus
REFERENCE
1 (bases 1 to 654)
AUTHORS
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Unpublished (1997)
JOURNAL
Tumor Gene Index
COMMENT
Contact: Robert Strausberg, Ph.D.

```

Email: cgapbs-remail.nih.gov
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
MGI:1397204
Seq primer: -40RP from Gibco
High quality sequence stop: 471.
Location/Qualifiers
1. 654

FEATURES

source

/organism="Mus musculus"
/db_xref="taxon:10090"
/clone="IMAGE:3514324"
/clone_lib="Soares_NKWM_mandible"
/tissue_type="mandible"
/lab_host="DH10B (phage-resistant)"
/note="Vector: p773D-Pac (Pharmacia) with a modified
polylinker; Site_1: NotI; Site_2: EcoRI; 1st strand cDNA
was primed with a Not I - oligo(dT) primer (5'
TGTTACCAATCTGAAGTGGAGCGCGCCCTTAATTTTTTTTTTTT 3'),
double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Not I and cloned into the Not I
and Eco RI sites of the modified p773 vector. Library
went through one round of normalization. Library
constructed by Bento Soares and M. Fatima Bonaldo."
BASE COUNT 228 a 118 c 137 g 168 t 3 others
ORIGIN

Query Match 18.9%; Score 644.6; DB 10; Length 654;
Best Local Similarity 98.9%; Pred. NO. 3e-100;
Matches 647; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 666 tttaaccccaatgatggaagaggaacacctcaacacattacttgggtccagttac 725
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DB 1 TTTTACCCCAATGATGATGGAAGAGGAACTCCACACATTACTTTGGTCCAGTAC 60
QY 726 tatttggcacagcattatgataaaattgggtcagccatccattgctctggaatacataa 785
|||||
DB 61 TATTTGGCACAGCATATGATAAAATTGGTCAGCCATCATTTGCTCGAATACATAAAT 120
QY 786 actgcaattgaaagtacacccaacttgatagaaactctttctgttaaaagctaaatctat 845
|||||
DB 121 ACTGCAATTGAAAGTACACCAACATTTGATAGAACTCTTTCTGTGTAAGCTTAAATCTAT 180
QY 846 aagcatgctgggaattattaaagaagctccagggtgattgattgaagccagccagccctggac 905
|||||
DB 181 AAGCATGCTGGGAATATTAAAGANGCTCAGGTGATGATGATGATGATGATGATGATGATGAT 240
QY 906 acagcagacagattattattcaatccaagtgctgcaaaatacatgtttataaaggccaacctgatt 965
|||||
DB 241 ACAGCAGACAGATTTATTATTTCAAGTGTGCAAAATACATGTTAAAAAGCCAACCTGATT 300
QY 966 aaagaggctgaagaaatgttccaagtttacgagggaaggaacttcagcggtgagaaac 1025
DB 301 AAAGAGGCTGAAGAAATGTTTCCAAAGTTTACGAGGGAAGAACTTCACGCGGTAGAGAAC 360
QY 1026 ctgaatgaatgcagtgattggttccagacagagtgctgcaggccatacaaaagcaatg 1085
DB 361 CTGAATGAATGCAGTGATGTTGTTCCACAGACAGATGCTGCTCAGGCATACAAAGCAATG 420
QY 1086 aacaaattggtgaagcaacttaagaatgtcatgaaattgagagacattttatagaatc 1145
DB 421 AACAAATTTGTTGAAGCACTTTAAGAAATGTCATGAAATTGAGAGACATTTATAGAAATC 480
QY 1146 accgatgaccagtttgactttcatatcatctgattgaggaagatcaccccttagatcatat 1205
DB 481 ACCGATGACCAAGTTGACTTTTCATACATACATGTTATGAGGAAGATCACCCCTTAGATCATAT 540
QY 1206 gtgacttattaaactagagaatgtactctgcagacagcatccattttacttcaaacgacgcg 1265
DB 541 GTGACTTATTAAACTAGAGATGTACTTCGACAGCATCCATTNTACTTCAAGCAGCG 600
QY 1266 agaattctattgagatctatttgaagcttctatgacacccctctgacagatgag 1319
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DB 501 AGAATTGCTATTGAGATCTATTGAAAGCTTCATGACACCCCTCTGCAGATGAG 654

RESULT 11
BB577716
LOCUS BB577716
DEFINITION BB577716 RIKEN full-length enriched, adult male medulla oblongata
MUS musculus cDNA clone 6330400115 5', mRNA sequence.
BB577716
ACCESSION BB577716
VERSION BB577716
KEYWORDS BB577716.2 GI:16449448
SOURCE EST.
ORGANISM house mouse.
MUS musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 673)
AUTHORS Arakawa,T., Carninci,P., Fukuda,S., Furuno,M., Hanagaki,T., Hara,A.,
, Hiramoto,K., Hori,F., Ishii,Y., Ito,M., Kawai,J., Konno,H., Kouda
, M., Koya,S., Matsuyama,T., Miyazaki,A., Nomura,K., Ohno,M.,
Okazaki,Y., Okido,T., Saito,R., Sakai,C., Sakai,K., Sano,H., Sasaki
, D., Shibata,K., Shinagawa,A., Shiraki,T., Sogabe,Y., Suzuki,H.,
Tagami,M., Tagawa,A., Takahashi,F., Takeda,Y., Tanaka,T., Toya,T.,
Muramatsu,M. and Hayashizaki,Y.
RIKEN Mouse ESTs (Arakawa,T., et al. 2001)
UNPUBLISHED (2001)
On Nov 30, 2000 this sequence version replaced gi:11474260.
Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-Ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@gs.c.riken.go.jp/
URL: http://genome.gsc.riken.go.jp/
Carninci,P., Shibata,Y., Hayatsu,N., Sugahara,Y., Shibata,K., Itoh
, M., Konno,H., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
Normalization and subtraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new
genes. Genome Res. 10 (10), 1617-1630 (2000)
wagi,K., Fujiwaki,S., Inoue,K., Togawa,Y., Izawa,M., Ohara,E.,
Watahiki,M., Yoneda,Y., Ishikawa,T., Ozawa,K., Tanaka,T., Matsuura
, S., Kawai,J., Okazaki,Y., Muramatsu,M., Inoue,Y., Kira,A. and
Hayashizaki,Y.
RIKEN integrated sequence analysis (RISA) system--384-format
sequencing pipeline with 384 multicapillary sequencer. Genome Res.
10 (11), 1757-1771 (2000)
Konno,H., Fukunishi,Y., Shibata,K., Itoh,M., Carninci,P., Sugahara
, Y. and Hayashizaki,Y.
Computer-based methods for the mouse full-length cDNA
encyclopedia: real-time sequence clustering for construction of a
nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
Kondo,S., Shinagawa,A., Saito,T., Kiyosawa,H., Yamanaka,I., Alzawa
, K., Fukuda,S., Hara,A., Itoh,M., Kawai,J., Shibata,K. and
Hayashizaki,Y.
Computational Analysis of Full-Length Mouse cDNAs Compared with
Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)
Please visit our web site (http://genome.gsc.riken.go.jp/) for
further details.
cDNA library was prepared and sequenced in Mouse Genome
Encyclopedia Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in RIKEN.
Division of Experimental Animal Research in Riken contributed to
prepare mouse tissues.
Location/Qualifiers
1. 673
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="6330400115"
/clone_lib="RIKEN full-length enriched, adult male medulla
oblongata"
/sex="male"

FEATURES
source

	QY	2752	gacacatc	2759		
	Ddb	661	GGACACATC	668		
RESULT	12					
LOCUS	AW260482/c					
DEFINITION	um80e10.x1 Sugano mouse liver mlia Mus musculus cDNA clone IMAGE:2317674 3' similar to WP:Y50D7_164.A CE22298 ; , mRNA sequence.					
ACCESSION	AW260482	GI:66333463				
VERSION	AW260482					
KEYWORDS	EST.					
SOURCE	house mouse.					
ORGANISM	Mus musculus					
Eukaryota; Metazoa;	Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 635)					
REFERENCE	Marra,M., Hillier,L., Kucaba,T., Martin,J., Beck,C., Wylie,T., Underwood,K., Steptoe,M., Theising,B., Allen.M., Bowers,Y., Person,B., Swaller,T., Gibbons,M., Pape,D., Harvey,N., Schurk,R., Ritter,E., Kohn,S., Shin,T., Jackson.Y., Cardenas,M., McCann.R., Waterston.R. and Wilson.R.					
AUTHORS	The WashU-NCI Mouse EST Project 1999					
TITLE	Unpublished (1999)					
JOURNAL	Other_ESTS: um80e10.v1					
COMMENT	Contact: Marra M/WashU-NCI Mouse EST project 1999 Washington University School of Medicine 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA Tel: 314 286 1800 Fax: 314 286 1810 Email: mouseest@wustl.edu This clone is available royalty-free through LLNL ; contact the IMAGE Consortium (info@image.llnl.gov) for further information. MG1:1010318 Seq primer: custom primer used High quality sequence stop: 508.					
FEATURES	Location/Qualifiers					
source	1..635 /organism="Mus musculus" /strain="C57BL" /db_xref="taxon:10090" /clone_image="IMAGE:2317674" /clone_lib="Sugano mouse liver mlia" /sex="female" /dev_stage="adult" /lab_host="DH10B" /notes="Organ: Liver; Vector: pME18S-FL3; Site:1: DraIII (CACTGTGTC); site:2: DraIII (CACCATGTG); 1st strand cDNA was primed with an oligo(dT) primer [ATGTGGCGCTTTTTTTTCTTTT] double-stranded cDNA was ligated to a DraIII adaptor [TGTTGGCTACTGC], digested and cloned into distinct draIII sites of the pME18S-FL3 vector (5' site CACTGTGTC, 3' site CACCATTGT). XhoI should be used to isolate the cDNA insert. Size selection was performed to exclude fragments <1.5kb. Library constructed by Dr. Sumio Sugano (University of Tokyo Institute of Medical Science). Custom primers for sequencing: 5' end primer CTCTGTCTCTAAAGCTGCC and 3' end primer CGACCTGCGACCTGCAGACA."					
BASE COUNT	152 a 137 c 111 g 235 t					
ORIGIN						
Query Match	18.3%; Score 623.8; DB 9; Length 635;					
Best Local Similarity	98.9%; Pred. No. 1.le-96;					
Matches	628; Conservative 0; Mismatches 7; Indels 0; Gaps 0					
QY	765	attgtcttggaatacaataactgcgaattgaaagtaccaccaattgatagaacctttt	824			
Dd	635	ATTGCTCTGGAATACATAAATACATGCAATTGAAAGTACCACCAATTATAGAACCTCT	576			

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QY 825 cttgttaaaagcctaaatctatacatgctgggaatattataaagaagctgcccaggtggatg 884
Db 575 CTTGTAAAGCCTACATCTATAGGATGCTGGGAATATTAAAGAGAGCTGCCAGGTGGATG 516
QY 885 gatgaagccagccctggacacagcagacagatttatttaattccaagtgtgcaaaatcac 944
Db 515 GATGAAGCCAGCCCTGGACACAGACAGACATTTATTAAATCCAAAGTGTGCAAAATAC 456
QY 945 atgttaaaagccaaactgattaaagagcctgaagaaatgtgtccaagtgttacaggggaa 1004
Db 455 ATGTTAAAGCCCAACTGATTAAAGAGCTGAAGAAATGTGTCCAAGTTTACGAGGAA 396
QY 1005 ggaactcagcggtgagaacctgaatgaatcgagtgatgtgttccagacagagtgt 1064
Db 395 GGAACCTTCTCGGTAGAGAACCTGAATGAATGCAGTGTGTGTGTTCCAGACAGAGTGT 336
QY 1065 gctcgggcatacaaaagcaatgaacaaatttggtgaagcacttaagaataatgcatagaatt 1124
Db 335 GCTCAGGCATACAAAGCAATGAACAAATTTGTTGAAGCACITTAAGAAATGTCATGAAT 276
QY 1125 gagagacattttatagaataaccagatgacacagtttgactttcattacatactgtatgag 1184
Db 275 GAGAGACATTTTATAGAATCACCAGTCCAGTGTGTGACTTTTCATACATACATGTTATGAG 216
QY 1185 aagatcccttagatcatatgtgacttataaaactagaagatgtacttcgacagcat 1244
Db 215 AAGATCACCTTAGATCATATGTGACCTTATTAACCTTAGAAGATGTACTTCGACAGCAT 156
QY 1245 ccattttacttcaagcgcgagaattgctattgagatctattgaaacttcacgacac 1304
Db 155 CCATTTTACTTCAAGACGCGAGGATTGCTATTGAGATCTATTGAAGCTTCATGACAA 96
QY 1305 cctctgacagatgagacaaagacacagcgtgacacagacacacacacacacacacacac 1364
Db 95 CCTCTGACAGATGAGACAAAGACACACAGGCTGATACAGCAACATGTAAGACAAAGAG 36
QY 1365 ctaagaabaactgcgtaataaacaagaagagctca 1399
Db 35 CTAAGAAGCTGCGGAATAAACAAGAAGAGCTCA 1

RESULT 13
BG080108
LOCUS
DEFINITION
H3049G08-5 NIA Mouse 15K cDNA Clone Set Mus musculus linear EST 26-JAN-2001
H3049G08 5', mRNA sequence.
ACCESSION
BG080108
VERSION
BG080108.1 GI:12562676
KEYWORDS
EST.
SOURCE
house mouse.
ORGANISM
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 625)
Kargul, G.J., Dudekula, D.B., Qian, Y., Lim, M.K., Jaradat, S.A., Tanaka
, T.S., Carter, M.G. and KO, M.S.H.
Verification and Initial annotation of NIA mouse 15K cDNA clone set
Unpublished (2001)
Other ESTs: H3049G08-3
Contact: George J. Kargul
Laboratory of Genetics
National Institute on Aging/National Institutes of Health
333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6820, USA
Email: cdna@igsun.grc.nia.nih.gov
This clone set has been freely distributed to the community. Please
visit http://igsun.grc.nia.nih.gov/cDNA/15k.html for details.
Plate: H3049 Row: G Column: 08
Seq primer: -21M13 Reverse
High quality sequence stop: 625
POLYA=NO.
FEATURES
Location/Qualifiers
source
1..625
/organism="Mus musculus"
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/strain="C57BL/6J"
/db_xref="niaEST:H3049G08-5"
/db_xref="taxon:10090"
/clone="H3049G08"
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/sex="Clones arrayed from a variety of cDNA libraries"
/dev_stage="Clones arrayed from a variety of cDNA
libraries"
/lab_host="DH10B"
/note="vector: pSPORT1; Site1: SalI; Site2: NotI; This
clone is among a rearrayed set of 15,247 clones from 11
embryo cDNA libraries (including preimplantation stage
embryos from unfertilized egg to blastocyst, embryonic
part of E7.5 embryos, extraembryonic part of E7.5 embryos
, and E12.5 female mesonephros/gonad) and one newborn
ovary cDNA library. Average insert size 1-5 kb. All
source libraries are cloned unidirectionally with Oligo(dT
)-Not primers. References include: (1) Genome-wide
expression profiling of mid-gestation placenta and embryo
using a 15,000 mouse developmental cDNA microarray, 2000,
Proc. Natl. Acad. Sci. U S A, 97: 9127-9132; (2)
Large-scale cDNA analysis reveals phased gene expression
patterns during preimplantation mouse development, 2000,
Development, 127: 1737-1749; (3) Genome-wide mapping of
unselected transcripts from extraembryonic tissue of
7.5-day mouse embryos reveals enrichment in the t-complex
and under-representation on the X chromosome, 1998, Hum
Mol Genet 7: 1967-1978."
BASE COUNT 211 a 111 c 143 g 160 t
ORIGIN

Query Match 18.2%; Score 623.4; DB 10; Length 625;
Best Local Similarity 99.8%; Pred. No. 1.2e-96;
Matches 624; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 422 aaaaatataggaagcctggactaaataccccaggggactcgtgccaagaaagctgcc 481
Db 1 AAAAATATATAGGAAGACCTGGACTAAATACCCAGGGGACTCGTGCACAGAGGCTGCC 60

QY 482 cttaaaactttttatctggagagaagtttaaggaggtgttgataggttctctaagagtgaa 541
Db 61 CTTAAACTTTTATCTCGAGAGAGTTTANGAGAGTGTGTGGATAGGTCTCTAAGGATCAA 120

QY 542 ttccagcaagggctgtccacctgtcttcaataccttgaggtctttatcacagagataaga 601
Db 121 TTTTCAGCAAGGGCTGTCCACCTGTCTCTCAATACCTTGAGGTCTTTATACAGAGATAAGA 180

QY 602 gaaggtggcaatcgtagaagaacttagttgattgaacacttcttaaaagtgtcg 661
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VERSION AI744486.1
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REFERENCE 1 (bases 1 to 710)
AUTHORS NCI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapsb-r@mail.nih.gov
Tissue procurement: Elias Campo, M.D., Michael R. Emmert-Buck, M.D.,
Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D.
cDNA Library Arraying: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CCAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
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was prepared from 12 pooled bulk tumor samples and primed
with a Not I - oligo(dT) primer. Double-stranded cDNA was
ligated to Eco RI adaptors (Pharmacia), digested with Not
I and cloned into the Not I and Eco RI sites of the
modified pT73 vector. Library went through one round of
normalization."
BASE COUNT 170 a 146 c 121 g 267 t 6 others
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VERSION AI130763.1
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SOURCE human.
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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 790)
AUTHORS Ota,T., Nishikawa,T., Suzuki,Y., Ishii,S., Saito,K., Kawai,Y.,
Yamamoto,J., Wakamatsu,A., Nakamura,Y., Nagai,T., Sugano,S. and
Isogai,T.
TITLE HRI human cDNA project
JOURNAL Unpublished (2000)
COMMENT Contact: Takao Isogai
Genomics Laboratory
Helix Research Institute
1532-3 Yana, Kisarazu, Chiba 292-0812, Japan
Tel: 81-438-52-3951
Fax: 81-438-52-3952
Email: genomics@hri.co.jp
HRI human cDNA project; 5'- & 3'-end one pass sequencing: Helix
Research Institute; cDNA library construction: Department of
Virology, Institute of Medical Science, University of Tokyo, and
Helix Research Institute.
FEATURES
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Job time: 14069 sec

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ORIGIN
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DB 61 AGTTGGCAACAACACTTGATGAATCTCTCCTAACAGAAACCTCCAGACATGTATGGAGG 120

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DB 710 GCACCTGGATTCTTTCACCTGAGCACAAAGAGTTGNTGGGCTTTAGCACTCTGACTGGATTT 769

QY 2659 gttacgggggttgggatt 2676
DB 770 GGTACNGGGNTGATCTTT 787
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us-09-836-410a-2.rst

Mon Jul 22 09:40:58 2002

OM of: US-09-836-410A-1 to: EST:* out_format : pfs

Date: Jul 20, 2002 3:07 AM

About: Results were produced by the GenCore software, version 4.5.
Copyright (c) 1993-2000 Compugen Ltd.

Command line parameters:

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-MINMATCH=0.100 -LOOPCL=0.000 -LOOPEXT=0.000 -QGAPOP=4.500
-QGAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500 -FGAPOP=6.000
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Search information block:

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Query length: 593

Database: EST*

Database sequences: 13736207

Database length: -1841457050

Search time (sec): 1584.610000

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gb_est2:BM0456794	1483.00	2224.26	1268	AGENCOURT_6404070 NIH
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gb_htc:AK007755	1374.00	2054.67	105	AK007755 Mus musculus 10 day c
gb_est2:BG067031	1180.00	1771.39	704	BM067031 H3049G08-3 NIA CGAP
gb_est1:AL744486	1176.00	1765.28	710	A1744486 w89h01.x1 NCI CGAP
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gb_est1:BM107262	1120.00	1681.61	652	BM107262 um56903.y1 Soares NKW
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gb_est2:BG080108	1093.00	1641.29	635	BM080108 H3049G08-5 NIA Mouse
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gb_est2:BF938468	904.00	1356.26	589	BF938468 fm77a09.y1 zebrafish
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DEFINITION Mus musculus adult male liver cDNA, RIKEN full-length enriched
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ACETYLTRANSFERASE, full insert sequence.
ACCESSION AK005056
VERSION AK005056.1 GI:12836717
KEYWORDS HTC; CAP trapper.
SOURCE Mus musculus (strain:C57BL/6J) adult male liver cDNA to mRNA,
clone:lib:RIKEN full-length enriched mouse cDNA library
clone:1300019C06.
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ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (sites)
AUTHORS Carninci, P. and Hayashizaki, Y.
TITLE High-efficiency full-length cDNA cloning
JOURNAL Meth. Enzymol. 303, 19-44 (1999)
MEDLINE 99279253
PUBMED 10349636
REFERENCE 2 (sites)
AUTHORS Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
TITLE Normalization and subtraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new genes
JOURNAL Genome Res. 10 (10), 1617-1630 (2000)
MEDLINE 20499374
PUBMED 11042159
REFERENCE 3 (sites)
AUTHORS Shibata, K., Itoh, M., Aizawa, K., Nagao, S., Sasaki, N., Carninci, P.,
Konno, H., Akiyama, J., Nishi, K., Katsunai, T., Tashiro, H., Itoh, M.,
Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Harada, A.,
Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K.,
Fujikawa, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M.,
Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsura, S., Kawai, J.,
Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
TITLE RIKEN integrated sequence analysis (RISA) system-384-format
sequencing pipeline with 384 multichannel sequencer
JOURNAL Genome Res. 10 (11), 1757-1771 (2000)
MEDLINE 20530913
PUBMED 11076861
REFERENCE 4 (sites)
AUTHORS The RIKEN Genome Exploration Research Group Phase II Team and the
FANTOM Consortium.
TITLE Functional annotation of a full-length mouse cDNA collection
JOURNAL Nature 409, 685-690 (2001)
MEDLINE 20530913
PUBMED 11076861
REFERENCE 5 (bases 1 to 2758)
AUTHORS Adachi, J., Aizawa, K., Akahira, S., Akimura, T., Aono, H., Arai, A.,
Arakawa, T., Baldarelli, R., Bono, H., Brownstein, M., Bul, C.,
Carninci, P., Fukuda, S., Fukunishi, Y., Furuno, M., Hanagaki, T.,
Hara, A., Hayatsu, N., Hill, D., Hiramoto, K., Hiraoka, T., Hori, F.,
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Numasaki, R., Ohno, M., Okazaki, Y., Okido, T., Owa, C., Quackenbush, J.,
Saito, H., Saito, R., Sakai, C., Sakai, K., Sano, H., Sasaki, D.,
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Sogabe, Y., Suzuki, H., Tagami, M., Tagawa, A., Takahashi, F.,
Tanaka, T., Tejima, Y., Toya, T., Yamamura, T., Yamanaka, I.,
Yasunishi, A., Yoshida, K., Yoshino, M., Muramatsu, M. and
Hayashizaki, Y.
```

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Direct Submission
Submitted (10-JUL-2000) Yoshihide Hayashizaki, The Institute of
Physical and Chemical Research (RIKEN), Laboratory for Genome
Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
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2436 AACTATAAAAAATATAATGTGAGAGCTCTAATAAGGTGTCTGAG 2485
536 euCysAspGlySerLeuArgAspCysLysGluAlaAlaGluAla 552
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2486 TCCTGGATGGCAGCTTGGGAAGTCTAGTCCCATATATGAGAGTA 2535
553 AlaSerCysHisLysLeuPheProTyrAlaLeuAlaPheMetPro 569
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2536 AGACCTGCCATAGCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 2585
569 yTyrGluGluAspMetLysIleThrValAsnGlyAspSerSer 586
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2586 G...AGGGAAGCCCTCGCTCTCAGTCTGAGTCTGAGTCTGAGT 2632
586 hrGluGluLeuAlaAsnGluIle 593
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2633 ACAAGCTCTTAATGAATTAAT 2655
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seq_name: gb_est2:BM456794

seq_documentation_block:

LOCUS BM456794

DEFINITION AGENCOURT_6404070 NIH_MGC_92 Homo sapiens cDNA clone IMAGE:5583707

1093 bp mRNA linear EST 05-FEB-2002

5', mRNA sequence.
BM456794
BM456794.1 GI:18505834

EST.
human.

ORGANISM
Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE
1 (bases 1 to 1093)

AUTHORS
NIH-MGC <http://mgs.nci.nih.gov/>.

TITLE
National Institutes of Health, Mammalian Gene Collection (MGC)

JOURNAL
Unpublished (1999)

COMMENT
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov

Tissue Procurement: ATCC

cDNA Library Preparation: Life Technologies, Inc.

cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

Cloning Strategy: Agencourt Bioscience Corporation

Cloning Distribution: MGC clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

<http://image.llnl.gov>

Plate: LLAM12347 row: d column: 12

High quality sequence stop: 695.

FEATURES
Location/Qualifiers

1..1093

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:5583707"

/clone_lib="NIH_MGC_92"

/tissue_type="embryonal carcinoma, cell line"

/lab_host="DH10B (phage-resistant)"

/note="Organ: testis; Vector: pCMV-SPORT6; Site_1: NotI;

Site_2: SalI; Cloned unidirectionally; oligo-dT primed.

Average insert size 2.5 kb. Library enriched for

full-length clones and constructed by Life Technologies.

Note: this is a NIH_MGC Library."

BASE COUNT 415 a 183 c 234 g 253 t 8 others

ORIGIN

alignment_scores:

Quality: 1483.00 Length: 336

Ratio: 4.678 Gaps: 8

Percent Similarity: 94.345 Percent Identity: 90.179

alignment_block:

US-09-836-410A-1 x BM456794 ..

Align seg 1/1 to: BM456794 from: 1 to: 1093

93 GlyLysGluGluProThrThrLeuLeuTrpValGlnTyrTyrLeuAl 109

2 GGAAGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 51

109 agLHisTyrAspLysLysLysLysLysLysLysLysLysLysL 126

52 ACAACATATGACAAATGCTGACGACATCTATTTGCTTTGGAGTACATA 101

126 snThrAlaIleGluSerThrProThrLeuLeuGluPheLeuValLys 142

102 ATACTGCTATTGAAAGTACACCTACATTAATAGAACTCTTTCTCGTCAA 151

143 AlaLysLysLysLysLysLysLysLysLysLysLysLysLysL 159

152 GCTAAAATCTATAAGCATGCTGGAATATTAAAGAGAGCTCAAGGTGGAT 201

159 tAspGluAlaGlnAlaLeuAspThrAlaAspArgPheIleAsnSerLysC 176

202 GGATGAGGCCAGGCTTGGACACAGACAGATTTATCACTCCAAAT 251

176 yAlaLysTyrMetLeuLysAlaAsnLeuLysLysLysLysLysL 192

252 GTGCAAAATACATGCTAAAAGCCCAACCTGATTAAAGAAAGCTGAAGAAATG 301

JOURNAL
COMMENT

Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cqbbs-r@mail.nih.gov
Tissue Procurement: DCTD/DTF
CDNA Library Preparation: Ling Hong/Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone Distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>

Plate: LHC1930 row: c column: 12
High quality sequence stop: 820.
Location/Qualifiers
1. 821

FEATURES

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/db_xref="taxon:9606"
/clone="IMAGE:5446979"
/clone_lib="NIH_MGC_40"
/tissue_type="carcinoma, cell line"
/lab_host="DH10B (phage-resistant)"
/note="Organ: prostate; Vector: pOB7; Site_1: XhoI;
Site_2: EcoRI; cDNA made by oligo-dT priming.
Directionally cloned into EcoRI/XhoI sites using the
following 5' adaptor: GGCACGAG(G). Library constructed by
Ling Hong in the laboratory of Gerald M. Rubin (University
of California, Berkeley) using ZAP-cDNA synthesis kit
(Stratagene) and Superscript II RT (Life Technologies).
Note: this is a NIH_MGC Library."

BASE COUNT 310 a 145 c 168 g 198 t
ORIGIN

alignment_scores:

Quality: 1401.00 Length: 274
Ratio: 5.151 Gaps: 2
Percent Similarity: 99.270 Percent Identity: 98.905

alignment_block:

US-09-836-410A-1 x BM044197

Align seg 1/1 to: BM044197 from: 1 to: 821

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2 CACAAAGAAAGAGTGGCAATCATAGAGAGTTAGTAGTTTAAAC 51
79 rSerLeuLysSerCysArgLeuPheAsnProAsnAspGlyLysGlu 96
52 CTCTCTAAAGAGTCCCGTTATTAAACCAATGATGGAAGAGG 101
96 LuProProThrThrLeuLeuTrpValGlnTyrTyrLeuAlaGlnHisTyr 112
102 AACCAACCAACACATTTACTTTGGTCCAGTACTTGGCACACATTTAT 151
113 AspLysIleGlyGlnProSerIleAlaLeuGluTyrIleAsnThrAla 129
152 CACAAATTTGGTCAGCCATCTATTCTTTGGAGTACATAAATACTGCTAT 201
129 eGluSerThrProThrLeuIleGluLeuPheLeuValLysAlaLysIle 146
202 TGAAGAGTACACCTACATTAATAGAACTCTTCTCGTGAAGCTAAATCT 251
146 YrLysHisAlaGlyAsnIleLysGluAlaAlaArgTrpMetAspGluAla 162
252 ATPAGCATGCTGGAATATTAAAGAGGTGCAAGGTGGATGGATGAGGCC 301
163 GlnAlaLeuAspThrAlaAspArgPheIleAsnSerLysCysAlaLysTy 179
302 CAGGCTTGGACACAGCAGACAGATTTATCAACTCCAAATGTGCAATA 351
179 rMetLeuLysAlaAsnLeuIleLysGluAlaGluMetCysSerLysP 196
352 CATGCTAAAGAACCACTGATTAAAGAAAGCTGAAGAAATGTGCTCAAGT 401

193 CysSerLysPheThrArgGluGlyThrSerAlaValGluAsnLeuAnG 209
302 TGCTCAAGATTTACAAGGGAAGAACATCAGCGGTAGAGAAATTTGAATGA 351
209 uMetGlnCysMetTrpPheGlnThrGluCysAlaGlnAlaTyrLysAla 226
352 AATGCAATGATGTTGTTCCAAACAGAAATGTCACAGGCTTTATTAAGCAA 401
226 eAsnLysPheGlyGluAlaLeuLysCysHisGluIleGluArgHis 242
402 TGAATAAATTTGGTGAACACTTAAGAAATGTCATGAGATTGAGAGACAT 451
243 PheIleGluLeuThrAspAspGlnPheAspPheHisThrTyrCysMetAr 259
452 TTTATAGAAATCACTGATGACCGCTTTGACTTTTCATACATGATGATGAG 501
259 gLysIleThrLeuArgSerTyrValAspLeuLeuLysLeuGluAspVal 276
502 GAAGATTACCTTATAGATCATATGTTGGACTTTATTAAGAACTAGAGATGTAC 551
276 euArgGlnHisProPheTyrPheLysAlaAlaArgIleAlaIleGluIle 292
552 TTCGACGACATCCATTTTACTTCAAGCAGCAGCAAGAAATGCTATAGATC 601
293 TyrLeuLysLeuHisAspAsnProLeuThrAspGluAsnLysGluHisG 309
602 TATTGAGGCTTCATGACACCCCTTACAGATGAGAAATAAAGAACACGA 651
309 uAlaAspThrAlaAsnMetSerAspLysGluLeuLysLysLeuArgAsn 326
652 AGCTGATACAGCAACATGCTCTGACAAAGAGCTAAAGAGCTACGTAATA 701
326 ysGlnArgArgAlaGlnLysLysAlaGlnIleGluGluGluLysLysAsn 342
702 KACAAAGAAAGAGCTCAAAAGAAAGCCAGATAGAGAGAGAAAGAAAT 751
343 AlaGluLysGluLysProGlnArgAsnProLysLysLysLysAspAspAs 359
752 GCCGAAAGAAAGAAAGCAGCAGCAAGAAATCAGAAAGAAAGAGGATGATGA 801
359 pAspGluGluIleGlyGlyProLysGluGluLeuIleProGluLysLeu 375
802 TGATGANGAGATAGGAGGCTCAAAAGAAAGAACTATTATCCAGAGAACTG 851
376 AlaLysValGluThrProLeuGluGluAlaIleLysPheLeuThrPr 391
852 GCCAAGGNTGAAAGCTCCATTTGGGAGAGAGCTATTAAATTTTAAACACC 901
391 oLeuLysAsn.....LeuValLysAsnLysIleGluThrHisLeuPhe 405
902 GGTGAAGAAAGCTGGTCAAGCAAGCAAGAAATAGAGGACCTCATCTTTT 951
406 Ala...PheGlu...IleTyrPheArgLysGluLysPheLeuLeuMet 419
952 TGCCCTTTTGAAGATTTACCTTTAAGGAAAGAAAGAAAGNTTCTCTCT 999

seq_name: gb_est2:BM044197

seq_documentation_block:
LOCUS BM044197 821 bp mRNA linear EST 07-NOV-2001
DEFINITION 603621442F1 NIH_MGC_40 Homo sapiens cDNA clone IMAGE:5446979 5',
mRNA sequence.
ACCESSION BM044197
VERSION BM044197.1 GI:16773464
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 821)
AUTHORS NIH-MGC <http://mgi.nci.nih.gov/>.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)

Mon Jul 22 09:40:55 2002

alignment_block:

US-09-836-410A-1 x AK007755

Align seg 1/1 to: AK007755 from: 1 to: 1959

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912 GAGAGGCTTCAGCTTTATGAGGAGTGCAGAACGAGCAGCACCACGAGAGT 961
19  uValProArgLysLeuProLeuAsnPheLeuSerGlyGluLysPheLysG 36
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
962 TTCACCCAGGAGGTCGCGTGAGTTTCGCCCCAGGTAAGAAGTTTCGAG 1011
36  LuCysLeuAspArgPheLeuArgMetAsnPheSerLysGlyCysProPro 52
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1012 AACTCATGATAAGTTCCTGAGACCAACTTTAGCAAAAGGTTGTCACCT 1061
53  ValPheAsnThrLeuArgSerLeuTyrArgAspLysGluLysValAlaI 69
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1062 CTGTTCACTACTTTGAAATCCTTGATGTGATGATACAGAAAAGGTTTCAAT 1111
69  eValGluGluLeuValGlyTyrGluThrSerLeuLysSerCysArgL 86
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1112 AATCCAGGAAGTGTGTTACTAATTATGAAGCCTCTCTTAAATGAATGGCT 1161
86  euPheAsnProAsnAspAspGlyLysGluGluProThrThrLeuLeu 102
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1162 APTTTAGGCTTTATGAGAACGGGGAAGAACCCCAACCACTCTAATC 1211
103  TrpValGlnTyrTyrLeuAlaGlnHisTyrAspLysIleGlyGlnProSe 119
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1212 TGGTTCAGTATTTCTTGGCAGCAGCATTTATGATAAATTTGGCAGTATT 1261
119  rIleAlaLeuGluTyrIleAsnThrAlaIleGluSerThrProThrLeuI 136
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1262 TCTGGCTTTGGAATATATTAATGCTGTAATTGCTAGTACTCAACTTAA 1311
136  leGluLeuPheLeuValLysAlaLysIleTyrLysHisAlaGlyAsnIle 152
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1312 TAGAACTATTTACATGAAAGCAAAANTTACACATATGGGTAAATC 1361
153  LysGluAlaAlaArgTrpMetAspGluAlaGlnAlaLeuAspThrAlaAs 169
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1362 AAAGAGCCGCACAGTGGATGATGAGGCACACAGTCTTTGGACACGGCTGA 1411
169  pArgPheIleAsnSerLysCysAlaLysTyrMetLeuLysAlaAsnLeuI 186
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1412 CAGGTTTCATCAATTCCAATGTGCCAAATGACATGCTTCGAGCAATATGA 1461
186  leLysGluAlaGluMetCysSerLysPheThrArgGluGlyThrSer 202
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1462 TAAAGACAGCAGGAAATGTGCTCCAGGTTTCAAGGGAAGGAACATCT 1511
203  AlaValGluAsnLeuAsnGluMetGlnCysMetTrpPheGlnThrGluCy 219
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1512 GCCATGGCAATCTGAATGAAATGCACTGTATGTGTTTGAGACGGAGTG 1561
219  sAlaGlnAlaTyrLysAlaMetAsnLysPheGlyGluAlaLeuLysLysC 236
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1562 CATTTTCGCTATCAGCGCTGGGAGATATGGGATGCTTGCTTGAATAAGT 1611
236  yHisGluIleGluArgHisPheIleGluIleThrAspAspGlnPheAsp 252
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1612 GCCATGAAGTAGAGGCACTTTCTTGAGATAACCGATGATCATGTTGAC 1661
253  PheHisThrTyrCysMetArgLysIleThrLeuArgSerTyrValAspLe 269
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1662 TTCCATACATCTACGATGAGAAAGATGACCTCCGTCCTTATGTTGCCT 1711
269  uLeuLysLeuGluAspValLeuArgGlnHisProPheTyrPheLysAlaA 286
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1712 CTTGAGATAGAAGATCTCTCAGAAAGACATACTTTTATTTTCAAGGCTG 1761
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286  iaArgIleAlaIleGluIleTyrLeuLysLeuHisAspAsnProLeuThr 302
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1762 CTAGATCAGCAATTGAAATATATTTGAAATTACATGATAACCTTTAACC 1811
303  AspGluAsnLysGluHisGluAlaAspThrAlaAsnMetSerAspLysG 319
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1812 AATGACAGCAAAACACAGACATAGATTTCAGAAAACCTGTACGCCAAGA 1861
319  uLeuLysLysLeuArgAsnLysGlnArgArgAlaGlnLysLysAlaGlnI 336
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1862 AATCAAGAAATGCTTAGCAAGCAAGAAAGAGCTCAGAAAAAGGCTAAGG 1911
336  leGluGluLysLysAsnAlaGluLysGluLysProGlnArgAsn 351
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1912 TAGAAGAAGAGAGAAAGCACACAGAAAGCGACGCAACACAGAAAAAC 1958
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seq_name: gb_est2.BG067031

seq_documentation_block: 704 bp mRNA linear EST 26-JAN-2001
LOCUS BG067031
DEFINITION H3049G08-3 NIA Mouse 15K cDNA Clone Set Mus musculus cDNA clone
H3049G08 3', mRNA sequence.

ACCESSION BG067031

VERSION BG067031.1 GI:12549600

KEYWORDS EST.

SOURCE house mouse.

ORGANISM

Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 704)

Kargul G.J., Dudekula,D.B., Qian,Y., Lim,M.K., Jaradat,S.A., Tanaka

,T.S., Carter,M.G. and Ko,M.S.H.

Verification and initial annotation of NIA mouse 15K cDNA clone set

Unpublished (2001)

Other ESTs: H3049G08-5

Contact: George J. Kargul

Laboratory of Genetics

National Institute on Aging/National Institutes of Health

333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6820, USA

Email: cdna@lgsun.grc.nia.nih.gov

This clone set has been freely distributed to the community. Please

visit <http://lgsun.grc.nia.nih.gov/cDNA/15k.html> for details.

Plate: H3049 row: G column: 08

Seq primer: -21M13 Forward

High quality sequence stop: 704

POLYA=Yes.

FEATURES

Location/Qualifiers

1..704

/organism="Mus musculus"

/strain="C57BL/6J"

/db_xref="niaEST:H3049G08-3"

/db_xref="taxon:10090"

/clone="H3049G08"

/clone_lib="NIA Mouse 15K cDNA Clone Set"

/sex="Clones arrayed from a variety of cDNA libraries"

/dev_stage="Clones arrayed from a variety of cDNA

libraries"

/lab_host="DH10B"

/note="Vector: pSPORT1; Site.1: SalI; Site.2: NotI; This

clone is among a rearrayed set of 15,247 clones from 11

embryo cDNA libraries (including preimplantation stage

embryos from unfertilized egg to blastocyst, embryonic

part of E7.5 embryos, extraembryonic part of E7.5 embryos

, and E12.5 female mesonephros/gonad) and one newborn

ovary cDNA library. Average insert size 1.5 kb. All

source libraries are cloned unidirectionally with Oligo(dT

)-Not primers. References include: (1) Genome-wide

expression profiling of mid-gestation placenta and embryo

using a 15,000 mouse developmental cDNA microarray, 2000,

Proc. Natl. Acad. Sci. U S A, 97: 9127-9132; (2)

Large-scale cDNA analysis reveals phased gene expression

patterns during preimplantation mouse development, 2000,

Development, 127: 1737-1749; (3) Genome-wide mapping of

unselected transcripts from extraembryonic tissue of 7.5-day mouse embryos reveals enrichment in the t-complex and under-representation on the X chromosome, 1998, Hum Mol Genet 7: 1967-1976."

BASE COUNT 168 a 149 c 124 g 263 t
ORIGIN

alignment_scores:

Quality: 1180.00 Length: 231
Ratio: 5.130 Gaps: 0
Percent Similarity: 99.567 Percent Identity: 99.567

alignment_block:

US-09-836-410A-1 x BG067031/rev ..

Align seg 1/1 to reverse of: BG067031 from: 1 to: 704

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692 GCACAGCATTTATGATAAAATTTGGTCAGCCATCCATTCTCTGGAATACAT 643
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125 eAsnThrAlaIleGluSerThrProThrLeuIleGluLeuPheLeuVal 142
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642 AAATCTGCAATTTGAAAGTACACCAACATTCATAGAACTCTTCTTGTA 593
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142 ysAlaLysIleTyrLysHisAlaGlyAsnIleLysGluAlaAlaAtgT 158
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592 AAGCTAAATCTATAAGCATGCTGGGAATATTAAAGNAGTCGCGGTGG 543
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159 MetAspGluAlaGlnAlaLeuAspThrAlaAspArgPheIleAsnSer 175
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542 ATGGATGAAGCCAGGCCCTGGACAGCAGACAGATTTTATTATTCCAA 493
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175 sCysAlaLysTyrMetLeuLysAlaAsnLeuIleLysGluAlaGluGlu 192
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492 GTGTGCAAAATACATGTTAAACCCCAACCTGTATTAAGAGCGCTGAAGA 443
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192 etCysSerLysPheThrArgGluGlyThrSerAlaValGluAsnLeuAs 208
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442 TGTGTTCCAAAGTTTACGAGGGAAGGAATTCACGCGTAGAAGACCTGA 393
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209 GluMetGlnCysMetTrpPheGlnThrGluCysAlaGlnAlaTyrLys 225
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392 GAATTCAGTGTATGTGGTCCAGACAGAGTGTGCTCAGGCATACAAAGC 343
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225 aMetAsnLysPheGlyGluAlaLeuLysLysCysHisGluIleGluArg 242
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342 AATGAACAAATTTGGTGAAGCACTTAAGAAATGTCATGAATTTGAGAC 293
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242 isPheIleGluIleThrAspAspGlnPheAspPheHisThrTyrCysMet 258
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292 ATTTTATAGAAATACCCGATGACCGATTTTGACTTTTCATACATCTGTAT 243
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259 ArgLysIleThrLeuArgSerTyrValAspLeuLeuLysLeuGluAspVa 275
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242 AGGAAGTACCCTTAGATCATATGTGGACTTATTAAAACTAGAAGATGT 193
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275 lleuArgGlnHisProPheTyrPheLysAlaAlaArgIleAlaIleGlu 292
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192 ACTTCGACAGCATCCATTTTACTTCAAAGCAGCGAGGATTCGATTGAGA 143
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292 leTyrLeuLysLeuHisAspAsnProLeuThrAspGluAsnLysGluHis 308
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142 TCTATTGAAAGCTTCATGACAAACCTCTGACAGATGAGAACAAAGAAC 93
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309 GluAlaAspThrAlaAsnMetSerAspLysGluLeuLysLysLeuArgAs 325
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92 GAGGCTGATACAGCAACATGTCTGACAAAGAGCTTAAAGAAATG.CGTA 44
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325 nLysGlnArgArgAlaGlnLysLysAlaGlnIleGluGlu 339
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43 TAAACAAAGAGAGCTCAAAAGAAAGCCAGCATAGAGAAGAG 1

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seq_documentation_block:

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ACCESSION: AI744486
VERSION: AI744486
KEYWORDS: EST.
SOURCE human.
ORGANISM Homo sapiens

REFERENCE

1 (bases 1 to 710)
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
NATIONAL Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index

JOURNAL:

Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgaps-r@mail.nih.gov
Tissue Procurement: Elias Campo, M.D., Michael R. Emmert-Buck, M.D., Ph.D.

COMMENT:

cDNA Library Preparation: M. Bento Soares, Ph.D.
cDNA Library Arraying: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: www-bio.llnl.gov/bbrp/image/image.html
Insert Length: 854 Std Error: 0.00
Seq Primer: -400P from GIBCO
High quality sequence stop: 462.

FEATURES

source

1..710
Location/Qualifiers
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/db_xref="taxon:9606"
/clone="IMAGE:2362801"
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/lab_host="DH10B"
/note="Vector: pT73D-Pac (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was prepared from 12 pooled bulk tumor samples and primed with a Not I - oligo(dT) primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT73 vector. Library went through one round of normalization."
BASE COUNT 170 a 146 c 121 g 267 t 6 others
ORIGIN

alignment_scores:

Quality: 1176.00 Length: 237
Ratio: 5.091 Gaps: 0
Percent Similarity: 97.468 Percent Identity: 96.203

alignment_block:

US-09-836-410A-1 x AI744486/rev ..

Align seg 1/1 to reverse of: AI744486 from: 1 to: 710

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710 TGGGTCAAGTACTTGGCACACCATATGACAAATTTGGTCAGCCATC 661
|||||
119 rleAlaLeuGluTyrIleAsnThrAlaIleGluSerThrProThrLeu 136
|||||
660 TANTGCTNTGGAGTACATAATACTGCTATTGANAGTACACT.ACATTAA 612
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136 leGluLeuPheLeuValLysAlaLysIleTyrLysHisAlaGlyAsnIle 152

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[illegible]

seq_name: gb_est2:BF012472

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seq_documentation_block:
LOCUS      BF012472                654 bp      mRNA      linear      EST 06-OCT-2000
DEFINITION ux56g03.y1 Soares_NKWMd_mandible mus musculus cDNA clone
IMAGE:3514324 5' similar to TR:Q9VW12 Q9VW12 CGI2202 PROTEIN. ; ,
mRNA sequence.
ACCESSION  BF012472
VERSION     BF012472.1  GI:10712747
KEYWORDS   EST.
SOURCE      house mouse.
ORGANISM   Mus musculus
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.; Mus.
            1 (bases 1 to 654)
REFERENCE  NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
AUTHORS   National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
TITLE      Tumor Gene Index
JOURNAL    Unpublished (1997)
COMMENT    Contact: Robert Strausberg, Ph.D.
            Email: cgapps-r@mail.nih.gov
            This clone is available royalty-free through LLNL ; contact the
            IMAGE Consortium (info@image.llnl.gov) for further information.
            MGI:11397204
Seq primer: -40RP from Gibco
            High quality sequence stop: 471.
FEATURES   Location/Qualifiers
            1..654
            /organism="Mus musculus"
            /db_xref="taxon:10090"
            /clone="IMAGE:3514324"
            /clone_lib="Soares_NKWMd_mandible"
            /tissue_type="mandible"
            /lab_host="DH10B (phage-resistant)"
            /note=vector: pYT30-Pac (Pharmacia) with a modified
            polylinker; Site_1: NotI; Site_2: EcoRI; 1st strand cDNA

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303 pGlu 304
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651 TGAG 654

seq_name: gb_est1:AW107262

seq_documentation_block:
LOCUS AW107262 652 bp mRNA linear 20-OCT-1999
DEFINITION um3c03.x1 Sugano mouse kidney mkia Mus musculus cDNA clone
IMAGE:2192164 3' similar to WP:Y50D7_164.A CE22298 ;, mRNA
sequence.
ACCESSION AW107262
VERSION AW107262.1 GI:6078062
KEYWORDS EST.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 652)
Marra, M., Hillier, L., Kucaba, T., Martin, J., Beck, C., Wylie, T.,
Underwood, K., Steptoe, M., Theising, B., Allen, M., Bowers, Y., Person
, B., Swaller, T., Gibbons, M., Pape, D., Harvey, N., Schurk, R., Ritter
, E., Kohn, S., Shin, T., Jackson, J., Cardenas, M., McCann, R.,
Waterston, R. and Wilson, R.
The WashU-NCI Mouse EST Project 1999
Unpublished (1999)
Contact: Marra M/WashU-NCI Mouse EST Project 1999
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1810
Email: mouseest@watson.wustl.edu
This clone is available royalty-free through LML; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
MGI:1004616
Seq primer: custom primer used
High quality sequence stop: 493.
Location/Qualifiers
1..652
/organism="Mus musculus"
/strain="C57BL"
/db_xref="taxon:10090"
/clone="IMAGE:2192164"
/clone_lib="Sugano mouse kidney mkia"
/sex="female"
/dev_stage="adult"
/lab_host="DH10B"
/notes="Organ: kidney; Vector: pME18S-FL3; Site_1: DraIII
(CACTGTG); Site_2: DraIII (CACCATGTG); 1st strand cDNA
was primed with an oligo(dT) primer
[ATGTGGCGCTTTTCTTTTCTTTT]; double-stranded cDNA was
ligated to a DraIII adaptor [TGTGGCGCTACTGG], digested
and cloned into distinct DraIII sites of the pME18S-FL3
vector (5' site CACTGTG, 3' site CACCATGTG). XhoI should
be used to isolate the cDNA insert. Size selection was
performed to exclude fragments <1.5kb. Library
constructed by Dr. Sumio Sugano (University of Tokyo
Institute of Medical Science). Custom primers for
sequencing: 5' end primer CTTCTGCTCTAAAAGCTGCG and 3' end
primer CGACCTGCAGCTCGAGCACA."
BASE COUNT 155 a 138 c 114 g 244 t 1 others
ORIGIN

alignment_scores:
Quality: 1120.00 Length: 217
Ratio: 5.185 Gaps: 0
Percent Similarity: 99.539 Percent Identity: 99.078

alignment_block:
US-09-836-410A-1 x AW107262/rev ..

Alignment seq 1/1 to reverse of: AW107262 from: 1 to: 652

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ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 625)
AUTHORS Kargul, G.J., Dudekula, D.B., Qian, Y., Lim, M.K., Jaradat, S.A., Tanaka, T.S., Carter, M.G. and Ko, M.S.H.
TITLE Verification and initial annotation of NIA mouse 15K cDNA clone set
JOURNAL Unpublished (2001)
COMMENT Other_ESTS: H3049G08-3
Contact: George J. Kargul
Laboratory of Genetics
National Institute on Aging/National Institutes of Health
333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6820, USA
Email: cdna@igsun.grc.nia.nih.gov
This clone set has been freely distributed to the community. Please
visit <http://igsun.grc.nia.nih.gov/cDNA/15k.html> for details.
Plate: H3049 row: G column: 08
Seq primer: -21M13 Reverse
High quality sequence stop: 625
POLYA-No.

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/db_xref="niaEST:H3049G08-5"
/db_xref="taxon:10090"
/clone="H3049G08"
/clone_lib="NIA Mouse 15K cDNA Clone Set"
/seq="Clones arrayed from a variety of cDNA libraries"
/db_stage="Clones arrayed from a variety of cDNA libraries"
/lab_host="DH10B"
/note="Vector: pSPORT1; Site 1: SalI; Site 2: NotI; This clone is among a rearranged set of 15,247 clones from 11 embryo cDNA libraries (including preimplantation stage embryos from unfertilized egg to blastocyst, embryonic part of E7.5 embryos, extraembryonic part of E7.5 embryos, and E12.5 female mesonephros/gonad) and one newborn ovary cDNA library. Average insert size 1.5 kb. All source libraries are cloned unidirectionally with Oligo(dT) -Not primers. References include: (1) Genome-wide expression profiling of mid-gestation placenta and embryo using a 15,000 mouse developmental cDNA microarray, 2000, Proc. Natl. Acad. Sci. U S A, 97: 9127-9132; (2) Large-scale cDNA analysis reveals phased gene expression patterns during preimplantation mouse development, 2000, Development, 127: 1737-1749; (3) Genome-wide mapping of unselected transcripts from extraembryonic tissue of 7.5-day mouse embryos reveals enrichment in the t-complex and under-representation on the X chromosome, 1998, Hum Mol Genet 7: 1967-1978."
Mol Genet 7: 1967-1978."

BASE COUNT 211 a 111 c 143 g 160 t
ORIGIN

alignment_scores:
Quality: 1093.00 Length: 208
Ratio: 5.255 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 99.519
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US-09-836-410A-1 x BG080108 ..
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2 AAAATATATAGAGAGCGCTGGACTAATACCCAGGGGACTCGTGCAAG 51
|||||
22 qLysLeuProLeuAsnPhelSerGlyGluLysPheLysGluCysLeuA 39
|||||
52 AAGGCTGCCCTTAACTTTTATCTCGAGAGAGATTAAAGGAGTGTGG 101
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39 spArgPheLeuArgMetAsnPhelSerLysGlyCysProProValPheAsn 55
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102 ATAGGTTCTTAAGATGAATTTTCAGCAAGGGCTGTCCACCTGCTTCAAT 151
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56 ThrLeuArgSerLeuTyrArgAspLysGluLysValAlaLeValGluGl 72
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152 ACCTTGAGGTCCTTATACAGAGATAAGAGAGGTGGCAATTCGTAGAAGA 201
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72 uLeuValValGlyTyrCluThrSerLeuLysSerCysArgLeuPheAsnP 89
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202 ACTAGTAGTTGGTTATGAAACTTCTCTAAAGAGTGTGCGCTATTAAAC 251
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89 roAsnAspAspGlyLysGluGluProProThrThrLeuLeuTrpValGln 105
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252 CCAATGATGATGGAAGAGGAGCACTCCAAACCACTTACTTTGGGTCCAG 301
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106 TyrTyrLeuAlaGlnHisTyrAspLysIleGlyGlnProSerIleAlaLe 122
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302 TACTATTGGCAGACAGCATTTATGATAAAATTTGGTCAGCCATCCATTG 351
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122 uGluTyrIleAsnThrAlaIleGluSerThrProThrLeuIleGluLeuP 139
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352 GGAATACATAAATACTGCAATTGAAAGTACACCAACATTTGATAGAACT 401
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139 heLeuValLysAlaLysIleTyrLysHisAlaGlyAsnIleLysGluAla 155
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402 TTCTTGTAAGCTAAATCTATAAGCATCTGGGAATATTAAAGAGCT 451
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156 AlaArgTrpMetAspGluAlaGlnAlaLeuAspThrAlaAspArgPheIl 172
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452 GCCAGGTGGATGGATGAAGCCAGCCCTGGACACAGACAGACATTTAT 501
|||||
172 eAsnSerLysCysAlaLysTyrMetLeuLysAlaAsnLeuIleLysGluA 189
|||||
502 TAATTCCAAGTGTCAAAATACATGTTAAAGCCCAACCTGATTAAAGAGG 551
|||||
189 iaGluGluMetCysSerLysPheThrArgGluGlyThrSerAlaValGlu 205
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552 CTGAAGAAATGTGTCCAAAGTTTACAGGGAAGGAACCTTCAGCGGTAGAG 601
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206 AsnLeuAsnGluMetGlnCysMet 213
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602 AACCTGAATGAATGCAGCTGTATG 625
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seq_name: gb_est1:AW260482
seq_documentation_block:
LOCUS AW260482 635 bp mRNA linear EST 23-DEC-1999
DEFINITION um80e10.x1 Sugano mouse liver mlia Mus musculus cDNA clone
IMAGE:2317674.3, similar to WP:Y50D7_164.A CE22298 ;, mRNA
sequence.
ACCESSION AW260482
VERSION AW260482.1 GI:66333463
KEYWORDS EST.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 635)
AUTHORS Marra, M., Hillier, L., Kucaba, T., Martin, J., Beck, C., Wylie, T., Underwood, K., Steptoe, M., Theising, B., Allen, M., Bowers, Y., Person, B., Swaller, T., Gibbons, M., Pape, D., Harvey, N., Schurk, R., Ritter, E., Kohn, S., Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R., and Wilson, R.
TITLE The WashU-NCI Mouse EST Project 1999
JOURNAL Unpublished (1999)
COMMENT Other_ESTS: um80e10.y1
Contact: Marra M/WashU-NCI Mouse EST Project 1999
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1810
Email: mouseest@watson.wustl.edu

REFERENCE:
AUTHORS.

1 (bases 1 to 988)
 Arakawa, T., Carninci, P., Fukuda, S., Furuno, M., Hanagaki, T., Hara, A., Hiramoto, K., Hori, F., Ishii, Y., Ito, M., Kawai, J., Konno, H., Kouda, M., Koya, S., Matsuyama, T., Miyazaki, A., Nomura, K., Ohno, M., Okazaki, Y., Okido, T., Saito, R., Sakai, C., Sakai, K., Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T., Sogabe, Y., Suzuki, H., Tagami, M., Tagawa, A., Takahashi, F., Takeda, Y., Tanaka, T., Toya, T., Muramatsu, M., and Hayashizaki, Y.
 RIKEN Mouse ESTs (Arakawa, T., et al. 2001)
 Unpublished (2001)
 On Nov 1, 1999 this sequence version replaced gi:6179369.
 Contact: Yoshihide Hayashizaki
 Laboratory for Genome Exploration Research Group, RIKEN Genomic
 Sciences Center (GSC), Yokohama Institute
 The Institute of Physical and Chemical Research (RIKEN)
 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
 Tel.: 81-45-503-9222

TITLE	JOURNAL	COMMENT
1. The Role of the Teacher in the Classroom	Journal of Educational Research	1980, Vol. 83, No. 1, pp. 1-10
2. The Impact of Technology on the Classroom	Journal of Educational Research	1980, Vol. 83, No. 2, pp. 11-20
3. The Role of the Student in the Classroom	Journal of Educational Research	1980, Vol. 83, No. 3, pp. 21-30
4. The Impact of the Teacher on the Student	Journal of Educational Research	1980, Vol. 83, No. 4, pp. 31-40
5. The Role of the Student in the Classroom	Journal of Educational Research	1980, Vol. 83, No. 5, pp. 41-50
6. The Impact of the Teacher on the Student	Journal of Educational Research	1980, Vol. 83, No. 6, pp. 51-60
7. The Role of the Student in the Classroom	Journal of Educational Research	1980, Vol. 83, No. 7, pp. 61-70
8. The Impact of the Teacher on the Student	Journal of Educational Research	1980, Vol. 83, No. 8, pp. 71-80
9. The Role of the Student in the Classroom	Journal of Educational Research	1980, Vol. 83, No. 9, pp. 81-90
10. The Impact of the Teacher on the Student	Journal of Educational Research	1980, Vol. 83, No. 10, pp. 91-100

RIKEN Mouse ESTs (Arakawa, T., et al. 2001)
Unpublished (2001)
On Nov 1, 1999 this sequence version replaced gi:6179369.
Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
tel: 81-45-503-9222

FEATURES
source

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Location/Qualifiers
1. .988
/organism="Mus musculus"
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/lab_host="DH10B"
/notes="Site 1: Sali; Site 2: BamHI; cDNA library was
prepared and sequenced in Mouse Genome Encyclopedia
Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in
RIKEN. Division of Experimental Animal Research in Riken
contributed to prepare mouse tissues. 1st strand cDNA wa
primed with a primer [5',
GAGAGAGAGAGATCCAGAGACTCTTTTTTTTTTTTTTTTNN 3'], cDNA wa
prepared by using trehalose thermo-activated reverse
transcriptase and subsequently enriched for full-length
cap-trapper. cDNA went through one round of normalization
to Rot = 10.0 and subtraction to Rot = 100.0. Second

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Ratio: 5.249
Percent Similarity: 98.558
Percent Identity: 98.558
Gaps: 0

alignment_block:
US-09-836-410A-1 x BE300741
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Align seg 1/1 to: BE300741 from: 1 to: 629

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3 AAAATCTATAAGCATGCTGCAATATATAAAGAAGCTCAAGGTTGGATGGA 52
160 pGluAlaGlnAlaLeuAspThrIlaAspArgPheIleAsnSerLysCysA 177
53 TGAAGGCCAGCGCTTGACACAGCAGACAGATTTATCACTCCAATCTG 102
177 laLysTyrMetLeuLysAlaAsnLeuIleLysGluAlaGluGluMetCys 193
103 CAAATACATGCTAAAGCCCAACCTGATTAAAGAAGCTGAAGAATGTGC 152
194 SerLysPheThrArgGluGlyThrSerAlaValGluAsnLeuAsnGluMe 210
153 TCAAAATTTACAAGGAAGGAACATCAGCGGTAGAGAATTTGAATCAAA 202
210 tGlnCysMetTrpPheGlnThrGluCysAlaGlnAlaTyrLysAlaMetA 227
203 GCAGTGCATGTGGTTCCAAACAGAAATGTGCCAGGCTTATAAAGCAATGA 252
227 snLysPheGlyGluAlaLeuLysLysCysHisGluIleGluArgHisPhe 243
253 ATAAATTTGGTGAACCACTTAAGAAATGTCATGAGATTGAGAGACATTT 302
244 IleGluIleThrAspAspGlnPheAspPheHisThrTyrCysMetArgLy 260
303 ATAGAAATCACTGATGACCAGTTTGACITTCATACACTCTGTATGAGGA 352
260 sileThrLeuArgSerTyrValAspLeuLeuLysLeuGluAspValIleuA 277
353 GATTACCCCTTAGCATCATATGTGCACCTTATTAAACTAGAAGATGTACTTC 402
277 rgGlnHisProPheTyrPheLysAlaAlaArgIleAlaIleGluIleTyr 293
403 GACAGCATCCATTTTACTTCAAGGCACAGAATTTGCTATAGAGATCTAT 452
294 LeuLysLeuHisAspAsnProLeuThrAspGluAsnLysGluHisGluAl 310
453 TTGAACCTTCATGACAAACCCCTTACAGATGAGAATAAAGAACAACGAAGC 502
310 aAspThrAlaAsnMetSerAspLysGluLeuLysLysLeuArgAsnLysG 327
503 TGTATACAGCAAAACATGCTGCACAAAGAGCTTAAGAAGCTACGTAATAAC 552
327 lnArgArgAlaGlnLysLysAlaGlnIleGluGluGluLysLysAsnAla 343
553 AAAGAGAGAGCTCANAAGAAAGGCCAGATAGAGAAGAGAGAAAAAATGCA 602
344 GluLysGluLysProGlnArgAsn 351
603 GAAAAGAAAAAGCAGCAGAGAAAT 626
acc name: gb_est1:AV227702

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seq_documentation_block: linear mRNA 988 bp EST 14-NOV-2001
 LOCUS AV227702 full-length enriched, 14 days embryo liver Mus
 DEFINITION AV227702 RIKEN full-length enriched, 14 days embryo liver Mus
 ACCESSION AV227702.1
 VERSION 1.0
 KEYWORDS
 SOURCE Mus musculus
 ORGANISM Mus musculus
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ACCESSION	AV227702	
VERSION	AV227702.2	GI:16385485
KEYWORDS	EST.	
SOURCE	house mouse.	
ORGANISM	Mus musculus	
	Eukaryota; Metazoa; Chordata; Mammalia; Eutheria; Rodentia	

SOURCE	ORGANISM	HOUSE MUSCULUS
	Mus musculus	
	Eukaryota; Metazoa;	Chordata; Vertebrata; Euteleostomi;
	Mammalia; Eutheria;	Muridae; Murinae; Mus.

Mon Jul 22 09:40:55 2002

us-09-836-410a-1.p2n.rst

Align seg 1/1 to: BJ057588 from: 1 to: 638

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173 nSerLysCysAlaLysTyrMetLeuLysAlaAsnLeuIleLysGluAlaG 190
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65 TTCTAAATGTCAAAATATATGTTAAAGCAACCTGATTAAAGAGGCTG 114
|||||
190 lbcGluMetCysSerLysPheThrArgGluGlyThrSerAlaValGluAsn 206
|||||
115 AAGAATGTGCTCGAAATTTACAGGAGGAGACATCAGCAGTGGAAAT 164
|||||
207 LeuAsnGluMetGlnCysMetTrpPheGlnThrGluCysAlaGlnAlaTy 223
|||||
165 CTGAACGAGATGCGATGTCATGTGGTCCAGACAGAAATGTGCACAAGCTTA 214
|||||
223 rLysAlaMetAsnLysPheGlyGluAlaLeuLysCysHisGluIleG 240
|||||
215 CAAATCCATGAATAATATGCGGAGGACCTTAAATAATGCCATGAAATG 264
|||||
240 luArgHisPheIleGluIleThrAspAspGlnPheAspPheHisThrTyr 256
|||||
265 AAAGGCATTTGTAGAAATAACAGATGACCAGTTTGATTTCCACACTTAC 314
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257 CysMetArgLysIleThrLeuArgSerTyrValAspLeuLysLeuG 273
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GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: July 19, 2002, 22:39:02 ; Search time 4131.3 Seconds
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Searched: 1797656 seqs, 10463268293 residues
Total number of hits satisfying chosen parameters: 3595312

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Post-processing: Minimum Match 0%
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Listing first 45 summaries

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- 33: em_htg_inv.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	3418	100.0	3418	6	AX285294	Sequence
3	3418	100.0	3418	6	AX285296	Sequence
4	3387	99.1	3421	10	AF237622	Sequence
5	2740.2	80.2	5505	9	HSA314788	Mus muscu
6	2543	74.4	4192	9	AF327722	Homo sapi
7	1779	52.0	1779	6	AX285247	Sequence
8	1469.4	43.0	3324	5	AF247679	Sequence
9	1413	41.3	1413	6	AX285295	Xenopus l
10	1210.2	35.4	1802	9	AK023387	Sequence
11	1191.8	34.9	101276	2	AC094440	Homo sapi
12	1126	32.9	2859	9	AK001595	Rattus no
13	1079.6	31.6	1985	9	AK023402	Homo sapi
14	1038.4	30.4	145395	9	AC097376	Homo sapi
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19	419	12.3	181573	2	AC020959	Mus muscu
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31	226	6.6	288	6	AX210600	Sequence
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ALIGNMENTS

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ACCESSION	AX285242.1	GI:17045930				
VERSION						
KEYWORDS						
SOURCE	human.					
ORGANISM	Homo sapiens					
REFERENCE	1 (sites)					
AUTHORS	Gendron,R.L. and Paradis,H.					
TITLE	Treatment of Ocular neovascularization and related diseases					
JOURNAL	Patent: WO 0179506-A 1 25-OCT-2001;					
FEATURES	Children's Hospital Research Foundation (US)					
source	Location/Qualifiers					
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121	cacagcagacatctctgataaagtggattatgaatatgaactcctcttatcaga	180			
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ACCESSION AX285294
VERSION AX285294.1 GI:17045975
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SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (sites)
AUTHORS Gendron,R.L. and Paradis,H.
TITLE Inhibition of bone tumor formation using antisense cdna therapy
JOURNAL Patent: WO 0179505-A 2 25-OCT-2001;
CHILDREN'S HOSPITAL MEDICAL CENTER (US)
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DEFINITION Sequence 4 from Patent WO0179505.
ACCESSION AX285296
VERSION AX285296.1 GI:17045977
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (sites)
AUTHORS Gendron,R.L. and Paradis,H.
TITLE Inhibition of bone tumor formation using antisense cdna therapy
JOURNAL Patent: WO 0179505-A 4 25-OCT-2001;
CHILDREN'S HOSPITAL MEDICAL CENTER (US)
FEATURES
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DEFINITION AF237622
ACCESSION AF237622
VERSION AF237622.1 GI:8164012
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Dev. Dyn. 218 (2), 300-315 (2000)
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2 (bases 1 to 3421)
Gendron, R.L., Adams, L.C. and Paradis, H.
Direct Submission
Submitted (20-FEB-2000) Pediatrics, Childrens Hospital Medical
Center, 3333 Burnet Avenue, Cincinnati, OH 45229-3039, USA
3 (bases 1 to 3421)
Gendron, R.L., Adams, L.C. and Paradis, H.
Direct Submission
Submitted (13-JUN-2000) Pediatrics, Childrens Hospital Medical
Center, 3333 Burnet Avenue, Cincinnati, OH 45229-3039, USA
REMARK Amino acid sequence updated by submitter
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VERSION AJ314788.1 GI:14589341
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SOURCE human.
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Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 5505)
AUTHORS Fluge O., Bruland O., Akslen, L.A., Varhaug, J.E. and Lillehaug, J.R.
TITLE Identification of NATH, a novel gene overexpressed in papillary
thyroid carcinomas
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 5505)
AUTHORS Fluge O.
TITLE Direct Submission
JOURNAL Submitted (31-MAY-2001) Fluge O., Dept. of Molecular Biology,
University of Bergen, Thormohlens gt 55, N-5020 Bergen, NORWAY
COMMENT related entry AF327722.
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DEFINITION Sequence 6 from Patent WO0179506.
ACCESSION AX285247
VERSION AX285247.1 GI:17045931
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (sites)
AUTHORS Gendron, R.L. and Paradis, H.
TITLE Treatment of ocular neovascularization and related diseases
JOURNAL Patent: WO 0179506-A 6 25-OCT-2001;
Children's Hospital Research Foundation (US)
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LOCUS			
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ACCESSION	AK023387		
VERSION	AK023387.1		
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SOURCE	Homo sapiens ovary, tumor tissue cDNA to mRNA, clone_lib:OVARC1		
ORGANISM	Homo sapiens		
REFERENCE	1 (sites)		
AUTHORS	Isogai,T., Ota,T., Hayashi,K., Sugiyama,T., Otsuki,T., Suzuki,Y., Nishikawa,T., Nagai,K., Sugano,S., Shiratori,A., Sudo,H., Wagatsuma,M., Hosoiri,T., Kaku,Y., Kodaira,H., Kondo,H., Sugawara,M., Takahashi,M., Chiba,Y., Ishida,S., Murakawa,K., Ono,Y., Takiguchi,S., Watanabe,S., Kimura,K., Murakami,K., Ishii,S., Kawai,Y., Saito,K., Yamamoto,J., Wakamatsu,A., Nakamura,Y., Nagahara,K., Masuo,Y., Ninomiya,K. and Iwayanagi,T.		
TITLE	NEO human cDNA sequencing project		
JOURNAL	Unpublished (2000)		
REFERENCE	2 (bases, 1 to 1802)		
AUTHORS	Isogai,T. and Otsuki,T.		
TITLE	Direct Submission		
JOURNAL	Submitted (23-AUG-2000) to the DBJ/EMBL/GenBank databases. Takao Isogai, Helix Research Institute, Genomics Laboratory; 1532-3 Yana, Kisarazu, Chiba 292-0812, Japan (E-mail:genomics@hri.co.jp, Tel:81-438-52-3951, Fax:81-438-52-3952)		
COMMENT	NEO human cDNA sequencing project supported by Ministry of International Trade and Industry of Japan; cDNA full insert sequencing: Research Association for Biotechnology; cDNA library construction, 5'- & 3'-end one pass sequencing and clone selection: Helix Research Institute (supported by Japan Key Technology Center etc.) and Department of Virology, Institute of Medical Science, University of Tokyo.		
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LOCUS Rattus norvegicus clone CH230-4F1, 101276 bp DNA linear HTG 20-DEC-2001
DEFINITION Rattus norvegicus clone CH230-4F1, *** SEQUENCING IN PROGRESS ***,
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ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
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51 unordered pieces.
AC094440
AC094440.2 GI:17941168
HTG; HTGS_PHASE1.
Norway rat.
Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus
1 (bases 1 to 101276)
Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-Osman,F.R., Allen,C.,
Alsbrooks,S.L., Amaratunge,H.C., Are,J.R., Banks,T., Barbarella,J.,
Benton,J., Blincke,K., Blankenbrow,K., Bonnin,D., Bouck,J.,
Bowle,S., Briele,M., Brown,E., Brown,M., Bryant,N.P., Buhay,C.,
Burch,P., Burkett,C., Burrell,K.L., Byrd,N.C., Carron,T.F.,
Carter,M., Cavazos,S., Chacko,J., Chacko,J., Chavez,D., Chen,G., Chen,R.,
Chen,Z., Chowdhry,I., Christopoulos,C., Cleveland,C.D., Cox,C.,
Coyle,M.D., Dederich,D.A., Delaney,K.R., Delgado,O.,
Denn,A.L., Ding,Y., Dinh,H.H., Douthwaite,K.J., Draper,H.,
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Elhaj,C., Escotto,M., Falls,T., Ferraguto,D., Flegg,N., Ford,J.,
Foster,P., Frantz,P., Gabisi,A., Gao,J., Garcia,A., Garner,T.,
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Hernandez,J., Hernandez,O., Hodgson,A., Hoques,M., Holloway,C.,
Hollins,B., Homs,F., Howard,S., Huber,J., Hulyk,S., Hume,J.,
Jackson,L.E., Jacobson,B., Jia,Y., Johnson,R., Jolivet,S.,
Joudah,S., Karlsson,E., Kelly,S., Khan,U., King,L., Korvah,J.,
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Nguyen,A., Nguyen,N., Nguyen,N., Nickerson,E., Nwokoko,S.,
Ogih,M., Okwunodu,G., Oragunye,N., Oviedo,R., Pace,A., Payton,B.,
Peery,J., Perez,L., Peters,L., Pickens,R., Primus,E., Pu,L.L.,
Quiles,M., Ren,Y., Rives,M., Rojas,A., Rojibokan,I., Rolfe,M.,
Ruiz,S., Savary,G., Scherer,S., Scott,G., Shen,H., Shoshitari,N.,
Slisson,I., Sodergren,E., Sonaite,T., Sparks,A., Stanley,H.,
Stone,H., Sutton,A., Svatek,A., Taber,P., Tamerisa,A., Tamerisa,K.,
Tang,H., Tansey,J., Taylor,C., Taylor,T., Telford,B., Thomas,N.,
Thomas,S., Usmani,K., Vasquez,L., Vera,V., Villalón,D., Vinson,R.,
Wall,R., Wang,S., Ward-Moore,S., Warren,R., Washington,C.,
Watlington,S., Williams,G., Williamson,A., Wleczyk,R., Wooden,S.,
Worley,K., Wu,C., Wu,Y., Wu,Y.F., Zhou,J., Zorrilla,S., Nelson,D.,
Weinstock,G. and Gibbs,R.
Direct Submission
Unpublished
2 (bases 1 to 101276)
Worley,K.C.
Direct Submission
Submitted (15-SEP-2001) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Dec 20, 2001 this sequence version replaced gi:15624274.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GAPL
Center clone name: CH230-4F1
----- Summary Statistics
Assembly program: Phrap; version 0.950329First call to
findPhrapList
Consensus quality: 81999 bases at least Q40
Consensus quality: 88613 bases at least Q30
Consensus quality: 93921 bases at least Q20
Estimated insert size: 77412; sum-of-contigs estimation
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Quality coverage: 0x in Q20 bases; agarose-fp estimation
 Quality coverage: 0x in Q20 bases; sum-Of-contigs estimation
 Quality coverage: 0x in Q20 bases; sum-Of-contigs estimation

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* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/genbank\_draft\_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 51 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

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[illegible]

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*	72108	73211:	contig	of 1104	bp in length
*	73212	73311:	gap	of unknown	length
*	73312	74467:	contig	of 1156	bp in length
*	73312	74567:	gap	of unknown	length
*	74468	75716:	contig	of 1149	bp in length
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*	76866	78104:	contig	of 1139	bp in length
*	76966	78204:	gap	of unknown	length
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*	78205	80469:	gap	of unknown	length
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*	80470	82300:	gap	of unknown	length
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*	82301	83933:	gap	of unknown	length
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*	85236	86801:	gap	of unknown	length
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*	93025	93124:	gap	of unknown	length
*	93125	94257:	contig	of 1133	bp in length
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*	94358	95684:	contig	of 1327	bp in length
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*	95685	96826:	contig	of 1042	bp in length
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*	98127	99641:	contig	of 1415	bp in length
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FEATURES source

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/db_xref="taxon:10116"
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Best Local Similarity 94.9%; Pred. No. 1.4e-228;
Matches 1367. Conservative 0; Mismatches 47; Indels 27; Gaps 12;

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VERSION AK023402.1 GI:10435324
KEYWORDS oligo capping; fjs (full insert sequence).
SOURCE Homo sapiens ovary, tumor tissue cDNA to mRNA, clone_lib:OVARC1
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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (sites)
AUTHORS Isogai,T., Ota,T., Hayashi,K., Sugiyama,T., Otsuki,T., Suzuki,Y.,
Nishikawa,T., Nagai,K., Sugano,S., Shiratori,A., Sudo,H.,
Wagatsuma,M., Hosoiri,T., Kaku,F., Kodaira,H., Kondo,H.,
Sugawara,M., Takahashi,M., Chiba,Y., Ishida,S., Murakawa,K.,
Ono,Y., Takiguchi,S., Watanabe,S., Kimura,K., Murakami,K.,
Ishii,S., Kawai,Y., Saito,K., Yamamoto,J., Wakamatsu,A.,
Nakamura,Y., Nagahari,K., Masuho,Y., Ninomiya,K. and Iwayanagi,T.
NEDO human cDNA sequencing project
Unpublished (2000)
REFERENCE 2 (bases 1 to 1985)
AUTHORS Isogai,T. and Otsuki,T.
TITLE Direct Submission
JOURNAL
REFERENCE 1
AUTHORS Isogai,T. and Otsuki,T.
TITLE Direct Submission
JOURNAL
COMMENT Genomics Laboratory; 1532-3 Yana, Kisarazu, Chiba 292-0812, Japan
(E-mail:genomics@nri.co.jp, Tel.81-438-52-3951, Fax:81-438-52-3952)
NEDO human cDNA sequencing project supported by Ministry of
International Trade and Industry of Japan; cDNA full insert
sequencing; Research Association for Biotechnology; cDNA library
construction, 5' & 3'-end one pass sequencing and clone selection;
Helix Research Institute (supported by Japan Key Technology Center
etc.) and Department of Virology, Institute of Medical Science,
University of Tokyo.
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McCarthy, M., McEwan, P., McKernan, K., McPheeters, R., Meldrim, J.,
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Norbu, C., Norman, C.H., O'Connor, T., O'Donnell, P., O'Neill, D.,
Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Pollara, V.,
Raymond, C., Retta, R., Rieback, M., Riley, R., Rise, C., Rogov, P.,
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Topham, K., Travers, M., Travis, N., Triglio, J., Vassiliev, H.,
Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.-J., Young, G.,
Zainoun, J., Zemsek, L., Zimmer, A. and Zody, M.
Direct Submission
Submitted (23-NOV-2001) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: <http://www-seq.wi.mit.edu>
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L19968
Center clone name: 511_H_12

* NOTE: This record contains 84 individual
* sequencing reads that have not been assembled into
* contigs. Runs of N are used to separate the reads
* and the order in which they appear is completely
* arbitrary. Low-pass sequence sampling is useful for
* identifying clones that may be gene-rich and allows
* overlap relationships among clones to be deduced.
* However, it should not be assumed that this clone
* will be sequenced to completion. In the event that
* the record is updated, the accession number will
* be preserved.
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* 668 767: gap of 100 bp
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41320 42018: contig of 699 bp in length
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42839 42938: gap of 100 bp
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TITLE
JOURNAL
COMMENT

* 43640 43739: gap of 100 bp
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Query Match 18.3%; Score 626.6; DB 2; Length 66729;
Best Local Similarity 98.7%; Pred. No. 1.6e-115;
Matches 684; Conservative 0; Mismatches 4; Indels 5; Gaps 5;

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Db 24493 TTTTCAATAACAGACCAGCTTCTTTTCTGCA 24525

Search completed: July 20, 2002, 02:41:30
Job time: 14548 sec

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: July 20, 2002, 00:31:22 ; Search time 356.49 Seconds
(without alignments)
16461.650 Million cell updates/sec

Title: US-09-836-410A-2

Perfect score: 3418

Sequence: 1 caagtaacaccgcgaagtg.....atgcaataaaattgtttggg 3418

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 1736436 seqs, 858457221 residues

Total number of hits satisfying chosen parameters: 3472872

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

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- 22: /SIDSI/gcgdata/hold-geneseq/geneseq-emb1/NA2001A.DAT:*
- 23: /SIDSI/gcgdata/hold-geneseq/geneseq-emb1/NA2001B.DAT:*
- 24: /SIDSI/gcgdata/hold-geneseq/geneseq-emb1/NA2002.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	3418	100.0	3418	24	AAD22687
2	3418	100.0	3418	24	Human tubedown-1 c
3	3418	100.0	3418	24	Human tubedown-1 b
4	1502.2	43.9	2477	23	DNA encoding novel
5	1413	41.3	2477	24	Human tubedown-1 b
6	1210.2	35.4	1802	22	Human cDNA sequenc
7	1126	32.9	2859	22	Human cDNA sequenc
8	1079.6	31.6	1985	22	Human cDNA sequenc
9	617.8	18.1	710	22	Human differential

10	616.2	18.0	790	22	AAH06489	Human cDNA clone (
11	565	16.5	1325	20	AAH99092	Human cancer cell
12	555.6	16.3	781	20	AAZ13705	Human gene express
13	555.6	16.3	781	20	AAH99053	Human validated ca
14	475.6	13.9	764	20	AAZ15983	Human gene express
15	475.6	13.9	764	20	AAH98777	Human validated ca
16	398.4	11.7	774	20	AAH98889	Human validated ca
17	343.6	10.1	488	22	AAH12222	Human cDNA clone (
18	342.6	10.0	408	22	AAH37727	Novel human diagno
19	340	9.9	408	22	AAH37726	Novel human diagno
20	339.8	9.9	402	22	AAH37350	Novel human diagno
21	310.6	9.1	773	20	AAH99063	Human validated ca
22	281	8.2	404	22	AAH66446	Human human polynu
23	276.6	8.1	727	22	AAH96842	Human neuroblastom
24	268	7.8	300	20	AAH98340	Human cancer cell
25	258.4	7.6	300	20	AAZ12696	Human gene express
26	258.4	7.6	300	20	AAH98342	Human cancer cell
27	242.6	7.1	399	20	AAH95559	Human cancer cell
28	239	7.0	255	22	AAH81986	EST clone CR392.
29	226	6.6	288	22	AAH81986	Rat differential t
30	215.4	6.3	255	22	AAH81733	Human differential t
31	214.4	6.3	300	20	AAH82031	Rat differential t
32	212.8	6.2	300	20	AAH98346	Human cancer cell
33	188	5.5	653	22	AAH07206	Human gene express
34	163.8	4.8	802	21	AAH02414	Human cDNA clone (
35	149.4	4.4	297	21	AAH21231	Human colon cancer
36	73.4	2.1	2703	10	AAH90541	Human secreted pro
37	73.4	2.1	2724	12	AAQ12226	DNA encoding N-alp
38	57.2	1.7	654	23	AAH69553	DNA encoding novel
39	57.2	1.7	654	23	AAH71133	DNA encoding novel
40	57.2	1.7	654	23	AAH75467	DNA encoding novel
41	57	1.6	450	22	AAH00120	Human reproductive
42	56	1.6	887	22	AAH94064	Human neuroblastom
43	55.6	1.6	14006	24	ABL33958	Human immune syste
44	54.2	1.6	6171	24	ABL32788	Human immune syste
45	53.8	1.6	12007	24	ABL32717	Human immune syste

ALIGNMENTS

RESULT 1

AAD22687
ID AAD22687 standard; CDNA; 3418 BP.
XX
AC AAD22687;
XX
DT 26-FEB-2002 (first entry)
XX
DE Tubedown-1 (tbdn-1) protein encoding CDNA.
XX

Tubedown-1 protein; tbdn-1; ophthalmological; cytostatic; vulnerary; cerebroprotective; angiogenesis inhibitor; ocular neovascularisation; retinal disease; diabetic retinopathy; retinopathy of prematurity; primary hyperplastic vitreous; macular degeneration; trauma; stroke; haemorrhagic shock; arthritis; arteriosclerosis; delayed wound healing; angiofibroma; granulation; nonunion fracture; retrolental fibroplasia; solid tumour growth; chronic glaucoma; sickle cell retinopathy; cancer; burn; scar; corneal neovascularisation; rubeosis iritis; uveitis; gene therapy; ss.

Unidentified.

XX	Key	Location/Qualifiers
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FT		/product= "Tubedown-1 protein #2"
FT	CDS	57..2189
FT		/*tag= b
FT		/product= "Tubedown-1 protein #3"
FT	CDS	87..2189
FT		/*tag= c
FT		/product= "Tubedown-1 protein #4"

FT	CDS		408..2189	
FT		/product=	d	
FT		*tag=	"tubedown-1 protein #1"	
FT	misc_feature	408..2186		
FT		/note=	"This region is specifically claimed as	
FT		SEQ ID NO: 6	in claim 5 of the specification"	
XX				
XX				
PN	WO200179506-A2.			
XX				
PD	25-OCT-2001.			
XX				
PF	17-APR-2001; 2001WO-US12548.			
XX				
PR	17-APR-2000; 2000US-197977P.			
XX	(CHIT-) CHILDRENS HOSPITAL RES FOUND.			
PA	Gendron RL, Paradis H;			
XX				
PI	WPI: 2002-026032/03.			
DR	P-PSDB; AAEL13589, AAEL13590, AAEL13591, AAEL13592.			
XX				
PT	Novel tubedown-1 protein comprising anti-angiogenic activity is useful			
PT	for treating angiogenesis-associated disease related to ocular			
PT	neovascularization, e.g., diabetic retinopathy, retinopathy of			
PT	prematurity -			
PS	Claim 10; Page 56-58; 85pp; English.			
XX	The present invention relates to tubedown-1 (tbdn-1) proteins and			
CC	their corresponding cDNAs. Tbdn-1 proteins having anti-angiogenic			
CC	activity are associated with acetyl transferase activity. They			
CC	regulate endothelial differentiation through protein acetylation,			
CC	DNA-binding or by interacting with and/or acetylating other protein			
CC	targets important for endothelial differentiation. In normal adult			
CC	eyes, tbdn-1 is expressed highly in the corneal endothelium proper			
CC	and in the vascular endothelium of the limbus and retina. Tbdn-1			
CC	proteins are useful for preventing, treating, inhibiting or delaying			
CC	the onset of angiogenesis-associated disease related to ocular			
CC	neovascularisation, e.g., a retinal disease, such as preferably			
CC	diabetic retinopathy or retinopathy of prematurity, or primary			
CC	hyperplastic vitreous, macular degeneration and any other conditions			
CC	involving ocular neovascularisation. Tbdn-1 proteins are also useful			
CC	for treating any pathological neovascularisation condition such as			
CC	head trauma, spinal trauma, stroke, haemorrhagic shock, arthritis,			
CC	arteriosclerosis, angiofibroma, delayed wound healing, granulations,			
CC	cancer, burns, scars, nonunion fractures, retrolental fibroplasia,			
CC	solid tumour growth. Proteins of the invention are also useful for			
CC	treating ocular neovascularisation conditions such as chronic glaucoma			
CC	sickle cell retinopathy, corneal neovascularisation, rubeosis iritis,			
CC	uveitis, neovascularisation of the optic nerve. Sequences of the			
CC	invention are also used in gene therapy. The present sequence is			
CC	a cDNA encoding tubedown-1 proteins.			
XX				
SQ	Sequence 3418 BP: 1157 A; 604 C; 704 G; 953 T; 0 other;			
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Best Local Similarity		100.0%;	Pred. NO. 0;	
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Oy	61	tttaccatttttagaagactatgaatggcgacaaataattttagaagadtttagaaaa	120	
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Oy	121	cacagcagacatctcgtataaagtggtattatgaatatagtggaactcctcttatcaga	180	
Db	121	cacagcagacatctcgtataaagtggtattatgaatatagtggaactcctcttatcaga	180	

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Db 3181 taccctcggaaacttttccacagtgctacaggtttgttaacttgaagccctcattctct 3240
Qy 3241 aagaataatttctcgtcagttgtttcaggcgaagcccaagactttgttaatttttaaaag 3300
Db 3241 aagaataatttctcgtcagttgtttcaggcgaagcccaagactttgttaatttttaaaag 3300
Qy 3301 gcccaagatttttttccaataacagacagcgtctcttttccctgcagttacaaatgtaat 3360
Db 3301 gcccaagatttttttccaataacagacagcgtctcttttccctgcagttacaaatgtaat 3360
Qy 3361 ttctttttttttttgtgttcaaaataaaggtaccaaatatgcaaatgtttttggg 3418
Db 3361 ttctttttttttttgtgttcaaaataaaggtaccaaatatgcaaatgtttttggg 3418

ID	AAH77156 standard; cDNA; 3418 BP.
XX	AAH77156;
AC	21-JAN-2002 (first entry)
DT	Human tubedown-1 cDNA.
XX	Human tubedown-1; tbdn-1; antisense; cytostatic; osteopathic;
DE	Human; tubedown-1; tbdn-1; antisense; cytostatic; osteopathic;
XX	tubedown-1; tbdn-1; antisense; cytostatic; osteopathic;
XX	bone tumour; osteosarcoma; Ewings sarcoma; metastasis; ss.
KW	Homo sapiens.
OS	Homo sapiens.
XX	Key Location/Qualifiers
FH	408..2189
FT	/tag= a
FT	/product= "tubedown-1"
FT	
XX	WO200179505-A2.
PX	25-OCT-2001.
XX	17-APR-2001; 2001WO-US12435.
PR	17-APR-2000; 2000US-197977P.
PR	17-APR-2001; 2001US-0836410.
XX	(CHIL-) CHILDRENS HOSPITAL RES FOUND.
PA	Gendron RL, Paradis H;
XX	WPI; 2002-017618/02.
PI	p-PSDB; AAG77907.
DR	Nucleic acid molecules antisense to the tubedown-1 gene prevent
XX	overexpression of tubedown-1 protein and are useful to treat
PT	osteosarcoma and Ewing's Sarcoma family of tumours
PT	Claim 1; Page 36-38; 56pp; English.
XX	The sequence represents a new human gene, tubedown-1 (tbdn-1). The
CC	invention relates to a novel isolated nucleic acid of the tubedown-1
CC	gene, and antisense nucleic acids to tbdn-1. The polynucleotides and
CC	protein of the invention have cytostatic and osteopathic activity. The
CC	polynucleotides of the invention may be used in antisense-therapy/gene
CC	therapy. They are useful in the treatment of bone tumours, especially
CC	osteosarcoma and Ewings sarcoma family of tumours. The compounds of the
CC	invention may also be useful for the prevention of metastases from these
CC	types of tumours, either alone or in combination with radiotherapy and/or
CC	chemotherapeutic agents.
XX	Sequence 3418 BP; 1157 A; 604 C; 704 G; 953 T; 0 other;
SQ	
Query Match	100.0%; Score 3418; DB 24; Length 3418;
Best Local Similarity	100.0%; Pred. No. 0;
Matches 3418; Conservative	0; Mismatches 0; Indels 0; Gaps 0;
Qy	1 caagtaacaccgcgaagatgatagagatctgacagtgagcatcgtattgttatgc 60
Dd	1 caagtaacaccgcgaagatgatagagatctgacagtgagcatcgtattgttatgc 60
Qy	61 ttaccattattagaactataaatggcagcaaaatttttagaagatttagaaaa 120
Dd	61 ttaccattattagaactataaatggcagcaaaatttttagaagatttagaaaa 120
Qy	121 cacagcacatctccgtataaagtgtattgaatatgtaactctcttatatcaga 180
Dd	121 cacagcacatctccgtataaagtgtattgaatatgtaactctcttatatcaga 180
Qy	181 atcaagttcttcggaagcaggcttttatagagaagccctggaacatctttgtacctatg 240
Dd	181 atcaagttcttcggaagcaggcttttatagagaagccctggaacatctttgtacctatg 240

QY 1321 acaagaacacgagcgtgacagcaaaacatgtctgacaaagagctaaagaaactcgcta 1380
Db 1321 acaagaacacgagcgtgacagcaaaacatgtctgacaaagagctaaagaaactcgcta 1380
QY 1381 ataaacaaagaagcgtcaaaaagaagccagattgaagaagagaaaaaaatgccgaaa 1440
Db 1381 ataaacaaagaagcgtcaaaaagaagccagattgaagaagagaaaaaaatgccgaaa 1440
QY 1441 aegaaagccgcgaacggaaatccgaaaaaagaaaaagagatgatgcgaagaatcgag 1500
Db 1441 aegaaagccgcgaacggaaatccgaaaaaagaaaaagagatgatgcgaagaatcgag 1500
QY 1501 gcccaaaagaagcgttatccctgagaactggccaaagtggccaaagtggaaactccatcggaag 1560
Db 1501 gcccaaaagaagcgttatccctgagaactggccaaagtggccaaagtggaaactccatcggaag 1560
QY 1561 ctattaaagtttttaacacacattgaagaacttggtagaagaacagatagaactcatcttt 1620
Db 1561 ctattaaagtttttaacacacattgaagaacttggtagaagaacagatagaactcatcttt 1620
QY 1621 ttgaccttgagatctacttttagaagaaaaagtttcttttgatgctacaactcagtaaaagc 1680
Db 1621 ttgaccttgagatctacttttagaagaaaaagtttcttttgatgctacaactcagtaaaagc 1680
QY 1681 gggcatttgcattgatttattagtcacccctgcttcagtgcatgctgactctttc 1740
Db 1681 gggcatttgcattgatttattagtcacccctgcttcagtgcatgctgactctttc 1740
QY 1741 attctgtgtgtaagtaagacttaccgaaacagagttagaacagattataaaacagaaa 1800
Db 1741 attctgtgtgtaagtaagacttaccgaaacagagttagaacagattataaaacagaaa 1800
QY 1801 tgaatgtcttttttgagcaacaaatcccaagaaattttaagtgaacacctttctgaaagga 1860
Db 1801 tgaatgtcttttttgagcaacaaatcccaagaaattttaagtgaacacctttctgaaagga 1860
QY 1861 attctgattcattgcacatagattatcagctgccaatggatattattattagattctt 1920
Db 1861 attctgattcattgcacatagattatcagctgccaatggatattattattagattctt 1920
QY 1921 ctagtcaaaaacagcaatagagctggcgacaacacttgatggatccctcccaacagaa 1980
Db 1921 ctagtcaaaaacagcaatagagctggcgacaacacttgatggatccctcccaacagaa 1980
QY 1981 accttcagacttgcaatggaagtgttggagccttgtgtgagctgagcctacagactgta 2040
Db 1981 accttcagacttgcaatggaagtgttggagccttgtgtgagctgagcctacagactgta 2040
QY 2041 aagaagctgcgaagcctcacagagcaagtgttcataaagctttcccttatgcttggctt 2100
Db 2041 aagaagctgcgaagcctcacagagcaagtgttcataaagctttcccttatgcttggctt 2100
QY 2101 tcatgctcctcctggatcaagagatagaagatcacagtgaacggagatagttctgcag 2160
Db 2101 tcatgctcctcctggatcaagagatagaagatcacagtgaacggagatagttctgcag 2160
QY 2161 aaacggaagaactggccaatgaaatctgaacatcattataaacaagcaaatggaatgacttt 2220
Db 2161 aaacggaagaactggccaatgaaatctgaacatcattataaacaagcaaatggaatgacttt 2220
QY 2221 ggaacatatctagtgataatatttttgcacgcacctgctgcaattgctcttacttacac 2280
Db 2221 ggaacatatctagtgataatatttttgcacgcacctgctgcaattgctcttacttacac 2280
QY 2281 agaatgagagagtaaatgttctgcttcaaaatagcttactacgttttttccctgctgaa 2340
Db 2281 agaatgagagagtaaatgttctgcttcaaaatagcttactacgttttttccctgctgaa 2340
QY 2341 aactatataaaatctaacattacagatagattaggttcagttctttaaataaaaa 2400
Db 2341 aactatataaaatctaacattacagatagattaggttcagttctttaaataaaaa 2400
QY 2401 gctgctaaaaattgagggttttaaaagaaaaaaaatccgctatctcttactctccct 2460

Db 2401 gctgctaaaaattgagggttttaaaagaaaaaaaatccgtaaccttactctccct 2460
QY 2461 tcccatgtttttaaactaaatttataataaactctggagcctataacacgttaacatacaggt 2520
Db 2461 tcccatgtttttaaactaaatttataataaactctggagcctataacacgttaacatacaggt 2520
QY 2521 gctggtgcagaaataataacttttaaaattgtcttctgtgagatcttctcagacagca 2580
Db 2521 gctggtgcagaaataataacttttaaaattgtcttctgtgagatcttctcagacagca 2580
QY 2581 taaaataaattgctgtttttagcactgattcttctcagacagcaatctcagacagca 2640
Db 2581 taaaataaattgctgtttttagcactgattcttctcagacagcaatctcagacagca 2640
QY 2641 tagcatctgcctgattcttctacgggttgggtgattgacataggaagatgcaatgca 2700
Db 2641 tagcatctgcctgattcttctacgggttgggtgattgacataggaagatgcaatgca 2700
QY 2701 atcactgtgtacagagcgtctacaacacatgcttgacatgcttgtagagactggacacata 2760
Db 2701 atcactgtgtacagagcgtctacaacacatgcttgacatgcttgtagagactggacacata 2760
QY 2761 gctacaaagcggattaaagtgaacccatagaaggtgttcaagtaacgtgtgtgttctccaaa 2820
Db 2761 gctacaaagcggattaaagtgaacccatagaaggtgttcaagtaacgtgtgtgttctccaaa 2820
QY 2821 attcactgtacatgatcagtttgggttcttctgtacacagtttttaaccggaagcaaccag 2880
Db 2821 attcactgtacatgatcagtttgggttcttctgtacacagtttttaaccggaagcaaccag 2880
QY 2881 ttggaacaattctcaatttaactaaacttgaaacttaaaataaactgcaaaactttat 2940
Db 2881 ttggaacaattctcaatttaactaaacttgaaacttaaaataaactgcaaaactttat 2940
QY 2941 catgttttggccaaaactgtttaaactgttaactgcaagaacccaaatgactgtgatg 3000
Db 2941 catgttttggccaaaactgtttaaactgttaactgcaagaacccaaatgactgtgatg 3000
QY 3001 caccacataattatgcaagcatgaattttcacctgagagtgaaaaaagaaactctacc 3060
Db 3001 caccacataattatgcaagcatgaattttcacctgagagtgaaaaaagaaactctacc 3060
QY 3061 atggcttgaagtacagagcagaactccctgactacacattctctatgactgtatgaagact 3120
Db 3061 atggcttgaagtacagagcagaactccctgactacacattctctatgactgtatgaagact 3120
QY 3121 aatatcaaaaacctcagcagccttcttcacgtatgcagataatgcagaaaaagtgctgctttaga 3180
Db 3121 aatatcaaaaacctcagcagccttcttcacgtatgcagataatgcagaaaaagtgctgctttaga 3180
QY 3181 taaccttgggaacttttccacagtgctacaggtttgttaaacttgaaacttgaaactcttctct 3240
Db 3181 taaccttgggaacttttccacagtgctacaggtttgttaaacttgaaacttgaaactcttctct 3240
QY 3241 aagaataataattctcgtcagttgttccagcgaagcccaagactttgttaatttttaag 3300
Db 3241 aagaataataattctcgtcagttgttccagcgaagcccaagactttgttaatttttaag 3300
QY 3301 gcccaagatttttttcaataacacagacagccttcttttccctgcagttacaaaatgtaat 3360
Db 3301 gcccaagatttttttcaataacacagacagccttcttttccctgcagttacaaaatgtaat 3360
QY 3361 ttctttttttttttgtgttcaaacataaagggtacacaaatgcaataaattgtttggg 3418
Db 3361 ttctttttttttttgtgttcaaacataaagggtacacaaatgcaataaattgtttggg 3418

RESULT 3
AAH77158/C
ID AAH77158 standard; cDNA; 3418 bp.
XX
AC AAH77158;

XX DT 21-JAN-2002 (first entry)
XX DE Human tubedown-1 base pairs 3418-1 antisense cDNA.
XX XX
KW Human; tubedown-1; tbdn-1; antisense; cytostatic; osteopathic;
KW bone tumour; osteosarcoma; Ewings sarcoma; metastasis; ss.
XX OS Homo sapiens.
XX OS
PN W0200179505-A2.
XX PD 25-OCT-2001.
XX PF 17-APR-2001; 2001WO-US12435.
XX XX
PR 17-APR-2000; 2000US-197977P.
PR 17-APR-2001; 2001US-0836410.
XX XX
PA (CHIL-) CHILDRENS HOSPITAL RES FOUND.
XX Gendron RL, Paradis H;
PI WPT; 2002-017618/02.
XX DR
XX PT Nucleic acid molecules antisense to the tubedown-1 gene prevent
XX overexpression of tubedown-1 protein and are useful to treat
XX osteosarcoma and Ewing's Sarcoma family of tumours -
XX XX
XX PS Claim 7; Page 39-41; 56pp; English.
XX XX
CC The sequence represents tubedown-1 (tbdn-1) bases 3418-1 antisense cDNA.
CC The invention relates to a novel isolated nucleic acid of the tubedown-1
CC gene, and antisense nucleic acids to tbdn-1. The polynucleotides and
CC protein of the invention have cytostatic and osteopathic activity. The
CC polynucleotides of the invention may be used in antisense-therapy/gene
CC therapy. They are useful in the treatment of bone tumours, especially
CC osteosarcoma and Ewings sarcoma family of tumours. The compounds of the
CC invention may also be useful for the prevention of metastases from these
CC types of tumours, either alone or in combination with radiotherapy and/or
CC chemotherapeutic agents.
XX XX
SQ Sequence 3418 BP; 953 A; 704 C; 604 G; 1157 T; 0 other;

Query Match 100.0%; Score 3418; DB 24; Length 3418;
Best Local Similarity 100.0%; Pred. No. 0;
Matches 3418; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 caagtaacacccgcaagatgatagaggatctgcagagtgcagcatcatgattggtatgc 60
DB 3418 CAAGTAACACCCCAAGATGATAGAGGATCTGCAGAGTGAGCATCATGATTGTTATGC 3359
QY 61 ttaccattattagaagactatgaatggcagcaaaaattttagaagagttttaggaaaa 120
DB 3358 TTACCATTATTAGAACACTATGAATGGCAGCAAAAATTTAGAGAGTTTAGAAAA 3299
QY 121 ccacagacatctcctgataaagtggaattgaatgaatagtaactcctctatatcaga 180
DB 3298 CACAGCAGACATCTCCTGATAAAGTGGATTTAGATATAGTGAATCTCCTTATATCAGA 3239
QY 181 atcaagttcttcgggaagcaggtcttttatagagaagcccttgaaacatttttgaactatg 240
DB 3238 ATCAAGTTCTTCGGGAAGCAGGCTTTATAGAGAAAGCCCTGGAACATCTTTGTACCTATG 3179
QY 241 aaaaagcagatttgatataaacttgctgttgaaagaacccaaaggaactctgttgcagt 300
DB 3178 AAAAGCAGATTCTGTATATAAATTCGTTGTTGAAGAACCAAGGGGAACCTCTGTTCAGT 3119
QY 301 tgtgtcgtttggaagatgctgctgacgttttatagagattacaagagaggaatcctgaaa 360
DB 3118 TGTCGCTTTGGNAGATGCTGCTGAGGTTTATAGAGGATTACAAGAGAGGAATCTCTGAAA 3059

QY 361 attggcctattacaaggcttagaaaaagcactgaagccagcactaatatgttagaacgac 420
DB 3058 ATTGGCCCTATTACAAAGCCTTAGAAAAGCACTGAAGCCAGCTAATATGTTAGAACGCG 2999
QY 421 taaaaatataggaagcctggactaaataccccaggggagactcgtgcagaagaaagctgc 480
DB 2998 TAAAAATATATGAGGAAGCCTGACTAAATACCCAGGGGACTCGTGCACAGAAAGCTGC 2939
QY 481 ccttaaaactttttatctggagagaagtttaagagagtgttttgataggttccctaaagatga 540
DB 2938 CCTTAAACTTTTATCTGGAGAGAGTTTAAAGAGTGTGGATAGGTTCCCTAAGATGA 2879
QY 541 atttcagcaaggcgtccacctgtcttcaataacctttgaggtcttttatcacagagataaag 600
DB 2878 ATTTCAGCAAGGGCTGTCCACCTGTCTTCAATACCTTGAGGTCTTTATACAGAGATAAAG 2819
QY 601 agaagtggaatcgttagaagaactagtagtttggttatgataaaactctctaaaaagtgtc 660
DB 2818 AGAAGGTGGCAATCGTAGAAGAACTAGTAGTTGGTTATGAAACTTCTCTAAAAGATTGTC 2759
QY 661 gctatttaaccccaatgatgtaggaagaggaacacctcaaccacattacttttgggtcc 720
DB 2758 GCCTATTTAACCCCAATGATGAGAAAGGAGGAACCTCCAACCCACATTACTTTGGGTCC 2699
QY 721 agtactatttggcacagcattatgataaaattgggtcagcgcacccattctctggaataca 780
DB 2698 AGTACTATTTTGGCACAGCAATTATGATAAAATTTGGTCAGCCATCCATTGCTCTGGAATACA 2639
QY 781 taaactactgcaattgaaagtacacacattgataaactcttctgttaaaagctcaaaa 840
DB 2638 TAAATACTGCAATTGAAAGTACACCAACATGTATAGAACTCTTTCTGTAAAAGCTAAAA 2579
QY 841 tctataagcatgctggtggaattattaaagaagctgccaggtggatggatgaagccagggccc 900
DB 2578 TCATTAAGCATGCTGGGAATATTAAAGAAGCTGCCAGGTGGATGGATGAAGCCAGGCC 2519
QY 901 tgaacacagcagacagatttatttaattccaaagtgtcaaaatacatgttaaaagccaacc 960
DB 2518 TGGACACAGCAGACAGATTATTAAATTCGAAGTGTGCAAAATACATGTTAAAAGCCAACC 2459
QY 961 tgattaaagagcgtgaagaatgtgttccaaagtttacagagggaaggaacttcagcggtag 1020
DB 2458 TGATTAAAGAGGCTGAAGAAATGTGTTCCAAGTTTACAGGGAAGGAACCTTCAGCGGTAG 2399
QY 1021 agaacctgaatgaatgcagtgatgtgtgttccagacagagtggtgcagcagcatacaag 1080
DB 2398 AGAACCTGAATGAATGCAGTGTATGTGGTTCACAGACAGAGTGTGCTCAGGCATACAAAG 2339
QY 1081 caatgaacaaatttgggtgaagcacttaagaatactgaataattgagagacattttatag 1140
DB 2338 CAATGAACAAATTTGGTGAAGCACCTTAAGAATGTGATGAATTTGAGAGACATTTTATAG 2279
QY 1141 aaatcacagatgaccagtttgactttcatatcactgtatgaggaagatacacccttagat 1200
DB 2278 AAATCACCGATGACCAGTTTGACTTTTCATACATCTGTATGAGGAAGATCACCCCTTAGAT 2219
QY 1201 catatgtggaacttataaaactagaagatgtaacttcacagacatccatttcaactcaag 1260
DB 2218 CATATGTGGACTTATTAAACTAGAAAGATGTACTTCGACACATCCATTTTACTTCAAAG 2159
QY 1261 cagcgaatgtctattgagatctatttgaagcttcacgaacccctctgacagatgaga 1320
DB 2158 CAGCGAATTTGCTATTGAGATCTATTGGAAGCTTCATGACAACCCCTCTGACAGATGAGA 2099
QY 1321 acaagaacacagaggtgatcacgaaacatgtctgacaaagagctaaagaaactgcta 1380
DB 2098 ACAAGAACACAGAGGCTCATACAGCAACATGTCTGACAAAAGAGCTAAGAAACTGCCGTA 2039
QY 1381 ataaacaagaagagctcaaaagaaagccagattgaagaagagaaaaaatgcccga 1440
DB 2038 ATAAACAAGAAGAGAGCTCAAAAGAAAGCCAGATTGGAAGAGAGAAAAAATGCCGAAA 1979
QY 1441 aagaaagccgcaagcgaatccgaaaaagaaaaagagatgatgatgacgaagaatgtgag 1500

XX WO200175067-A2.
 XX 11-OCT-2001.
 XX 30-MAR-2001; 2001WO-US08631.
 XX 31-MAR-2000; 2000US-0540217.
 XX 23-AUG-2000; 2000US-0649167.
 XX (HYSE-) HYSEQ INC.
 XX Drmanac RT, Liu C, Tang YT;
 XX WPI: 2001-639362/73.
 XX P-PSDB; ABG07738.
 XX New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -
 XX
 PS Claim 1; SEQ ID NO 7729; 103pp; English.
 XX The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AA564197-AA594564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 XX
 XX Sequence 2477 BP; 859 A; 437 C; 522 G; 659 T; 0 other;
 XX
 Query Match 43.9%; Score 1502.2; DB 23; Length 2477;
 Best Local Similarity 80.7%; Pred. No. 0;
 Matches 2082; Conservative 0; Mismatches 218; Indels 279; Gaps 17;
 QY 27 gatctgcagatgagcatcatggttggtatgct-----ttacattattagaagact 81
 DB 149 gacctgcgacagacatcatggttggtatgcttattgcttaccattattagaagatt 208
 QY 82 atgaattggcagcaaaatttagaagatttagaagaacacacagacacatctctgata 141
 DB 209 atgaattggcagcaaaatttagaagatttagaagaacacacacagacacatctctgata 268
 QY 142 aagtgattatgaatagtagtaacctctcttatatcagaatacaagttcttcggaagcag 201
 DB 269 aagtgattatgaatagtagtaacctctcttatatcagaatacaagttcttcggaagcag 328
 QY 202 gtctttatagaagcccttggaacatctttgtacattatgaagaagcagatttgtataaac 261
 DB 329 gtctctatagaagccttttggaacatctttgtacattatgaagaagcagatttgtataaac 388
 QY 262 ttgctgttgaagaacaaagggaactctgttgcagttgtgtcgtttggaagatgctg 321
 DB 389 ttgctgttgaagaacaaagggaactctgttgcagttgtgtcgtttggaagatgctg 448

QY 322 ctgacgtttatagaggattacaagagagggaatcctgaaattgggcttattacaaggct 381
 DB 449 -cagatgtttatagaggattgcaagagagaatcctgaaactggcctattacaaggct 508
 QY 382 tagaaaaagcactgaagccagcctaattgttagaacggcctaaaaatatatgaggaagcct 441
 DB 509 tggaaaaagcactgaagccagcctaattgttagaacggcctaaaaatatatgaggaagcct 568
 QY 442 ggactaaataccacaggggactcgtccaagaagagctgccttaaaactttttatctggag 501
 DB 569 ggactaaataccacaggggactcgtccaagaagagctgccttaaaactttttatctggag 628
 QY 502 agaagtttaagagtggttgataggttcttaagatgaatttcagcaagggtgtctcac 561
 DB 629 agaagtttaagagtggttgataggttcttaagatgaatttcagcaagggtgtctcac 688
 QY 562 ctgttccaatactctgaggtcttta-tacagagataagagaag-gtggcaactgtaga 619
 DB 689 cagcttccaatactttaagatcattactaccaagacaagaagagtggtgcaactataga 748
 QY 620 agaactagtagtggttatgaaactctctaaaaagttgtgccttatttaaccccaatga 679
 DB 749 aaagttagtagtagtggttatgaaactctctaaaaagctgcgcgttatttaaccccaatga 808
 QY 680 tgatgaaaggaggaaacctccaaccacattactttgggtccagttactatttggcacagca 739
 DB 809 tgatgaaaggaggaaacctccaaccacattactttgggtccagttactatttggcacagca 868
 QY 740 ttatgataaaattgctcagccatccattgctggaatacataataatactgcgaattgaaag 799
 DB 869 ttatgataaaattgctcagccatccattgctggaatacataataatactgcgaattgaaag 928
 QY 800 tacaccaactgatagaactcttttctgaaaaagcctaaactctataaagcagctgggaa 859
 DB 929 tacacctataattagaactctttctctgaaaagcctaaactctataaagcagctgggaa 988
 QY 860 tattaaagagctgcagtgatgagatgaagccagccctgggacacagcagacagatt 919
 DB 989 tattaaagagctgcagtgatgagatgaagccagccctgggacacagcagacagatt 1048
 QY 920 tattaattccaagtgtgcaaaatcacatgtttaaagcccaacctgatttaaagagagctgaa 979
 DB 1049 tatcaactcccaatgtgcaaaatcacatgtttaaagcccaacctgatttaaagagagctgaa 1108
 QY 980 aatgtgtccaagtgttacgaggaagaaacttcagcgttagagaacctgaataaagcga 1039
 DB 1109 catgagctcaagtttacaaaggggacacatcagcgttagagaatttgaaatgaattca 1168
 QY 1040 gtatgtgtgttcacagacagatgtgctcaggcatacaagcaatgaacaaatttgggtga 1099
 DB 1169 gtcatgtgtgttcacatacagaagtgtcccagcttataaagcaatgaataatttgggtga 1228
 QY 1100 agcacttaagaatgtcatgaaattgagagacattttatagaataccagatcacagctt 1159
 DB 1229 agcacttcagaatgtcatgagattgagagacattttatagaataccagatcacagctt 1288
 QY 1160 tgactttcatatactactgtatgaggaagatcacaccttagatcatatgtggaactattaaa 1219
 DB 1289 tgactttcatatactactgtatgaggaagatcacaccttagatcatatgtggaactattaaa 1348
 QY 1220 actagaagatgtacttcgacagcatccattttacttctcaag--cagcgagaattgctatt 1277
 DB 1349 actagaagatgtacttcgacagcatccattttacttcaagggcgagcaagaattgctata 1408
 QY 1278 gagatctattttaa-gcttcatgacaacctctgacagatgagacaagaacacagcggc 1336
 DB 1409 gagatctatttgaaggcttcatgacaacctctgacagatgagacaagaacacagcggc 1468
 QY 1337 tg-----atacagcaaacatgtctgacaagagctcaaaagactgcaataaacaag 1390
 DB 1469 aggtcggtacagcccaaacatgtctgacaagagctcaaaagagctcaataaacaag 1528
 QY 1391 aagagctcaaaagaaagcccgatttgaagaagagaaaaaaatccgaaaaaagcc 1450

181 atcaagttcttcggaagcaggtctttatagagaagccctggaaacatctttgtacattg 240
1233 ATCAAGTTCTTCGGAAGCAGGCTTTATAGAGAAGCCTTGAACATCTTTGTACCTATG 1174
241 aaagcagatttgataaacttctgttgagaacacaaagggaactcttctgttcagt 300
1173 AAAAGCAGATTGTGATAAATCTCTGTGTGAAGAACCAAGGGGAACCTCTGTTCAGT 1114
301 tctgtcgtttgaaagctgctgacgtttatagaggattacaagagaggaatcctgaaa 360
1113 TGTGTCGTTGGAAGATCTGCTGACGTTTATAGAGATTACAAGAGAGGAATCCTGAAA 1054
361 attggccattacaaagccttagaanaagcactaaagcagctaatgttagaacgac 420
1053 ATTGGGCTTATTACAAAGCCTTAGAAAAGCCTGAAGCAGCTTAATATGTAGAACGCG 994
421 taaaaatataggaagcctgactgaactaaatccccagggactgctgcaagaagcgtgc 480
993 TAAAAATATAGGAGAGCCTGGACTAAATACCCAGGGGACTCGTGGCAAGAAAGCTGC 934
481 ccttaacttttatctgagagaaggttaagagaggttttgataggttcttaagatga 540
933 CCTTAACTTTTATCTGAGAGAGATTAAAGAGGTGTTGGATAGGTTCTTAAGATGA 874
541 atttcaagagggctgtccacctgtctcaatccttgaggtctttatcacagagataag 600
873 ATTTACAGCAAGGGCTGTCCACCTGCTCTCAATACCTTGAGGTTTATACAGAGATAAG 814
601 agaagtggaactcgttagaagaactagttggtttatgaactcttcaaaaagtgtgc 660
813 AGAAGTGGAATCGTAGAAGAACTAGTAGTTGTTATGAAACTTCTTAAAGAGTTGTC 754
661 gctatttaaccccaatgatagaaagaggaagcctcaacacatctacttgggtcc 720
753 GCCTATTAAACCCCAATGATGATGGAAGAGGAGAACTCCCAACACCATTTACTTTGGTCC 694
721 agtactattggcacagcattatgaataaattggtcagccatccattgctctggaataa 780
693 AGTACTATTGGCCACACATATGATAAATTTGTCAGCCATCCATTGCTCTGGAATACA 634
781 taaactatgcaattgaagtacacacacattgatagaactctttctgttaaaagctaaa 840
633 TAAATCTGCAATTGAAAGTACACCAACATTTGATAGAACTCTTTCTTGTAAAGCTAAA 574
841 tctataagcagctggaataataaaagctgcaagtggtgatgagagccagccccc 900
573 TCTATAGCATGCTGGGAATATTAAAGATGATGAGTGCAGGTCGATGATGAAGCCAGGCC 514
901 tggcacagcagacagatttattattccaaagtgtgcaaaatacatgtttaaagccaac 960
513 TGGACACAGCAGACAGATTTTATTATTCCTCAAGTGTGCAAAATACATGTTAAAGCCCAAC 454
961 tgattaaagagcctgaagaatgtttccaaagtttacagaggaaggaacttcagcgtag 1020
453 TGAATTAAGAGGCTGAAGAAATGTGTTCCAAAGTTTACGAGGAGGAACCTTCACGGTAG 394
1021 aqaacctaaatgaatgacgt 1080
393 AGAAGCTGAATGAATGACGATGTATGTGTTCCAGACAGAGTGTCTCAGGCATACAAAG 334
1081 caatgaacaaatttggtagaaccttaagaatgtcatgaaattgagagacattttatag 1140
333 CAATGAACAAATTTGGTGAAGCACTTAAGAAATGTGCAAAATACATGTTAAAGCCCAAC 274
1141 aaatcacagatgacaggtttgactttacatactgtatgaggaagatcaccttagat 1200
273 AAATCACCATGACAGGTTGACTTTTATACATATCTGTATGAGGAGATCACCTTAGAT 214
1201 catatgtgactattaaacttagaagatgtacttcgcagagatccattttacttcaag 1260
213 CATATGTGGACTATTAAACCTAGAGATGTCTTCGACAGCATCCATTTTACTTCAAG 154
1261 cagcagaattgctattgagatctatttgaagcttcatgacaacccctctgacagatgaga 1320

153 CAGCGAATGCTATTTGAGATCTATTTGAAGCTTCATGACAACCCCTCTGACAGATGAGA 94
1321 acaagaacacagcaggtgatcacagcaaacatgtctgtacaaagagctaaagaaactcgta 1380
93 ACAAGAACAACGAGGCTGTATACAGCAAAACATGCTGTGACAAGAGCTTAAGAAACTGCGTA 34
1381 ataacaagaagagctcaaaagaaagcccgaga 1413
33 ATAAACAAGAAGAGCTCAAAAGAAAGGCCGAGA 1
RESULT: 6
AAH16408
ID AAH16408 standard; cDNA; 1802 BP.
XX AC AAH16408;
XX XX 26-JUN-2001 (first entry)
XX XX Human cDNA sequence SEQ ID NO:15380.
XX DE Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.
XX KW Homo sapiens.
XX OS Homo sapiens.
XX PN EPI074617-A2.
XX XX 07-FEB-2001.
PD 28-JUL-2000; 2000EP-0116126.
PF 29-JUL-1999; 99JP-0248036.
XX PR 27-AUG-1999; 99JP-0300253.
PR 11-JAN-2000; 2000JP-0118776.
PR 02-MAY-2000; 2000JP-0183767.
PR 09-JUN-2000; 2000JP-0241899.
XX XX (HELI-) HELIX RES INST.
XX XX Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX WPI; 2001-318749/34.
XX XX Claim 8; SEQ ID 15380; 2537pp + CD ROM; English.
XX CC The present invention describes primer sets for synthesizing 5602
CC full-length cDNAs defined in the specification. Where a primer set
CC comprises: (a) an oligo-dT primer and an oligonucleotide complementary
CC to the complementary strand of a polynucleotide which comprises one of
CC the 5602 nucleotide sequences defined in the specification, where the
CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination
CC of an oligonucleotide comprising a sequence complementary to the
CC complementary strand of a polynucleotide which comprises a 5'-end
CC sequence and an oligonucleotide comprising a sequence complementary to a
CC polynucleotide which comprises a 3'-end sequence, where the
CC oligonucleotide comprises at least 15 nucleotides and the combination of
CC the 5'-end sequence/3'-end sequence is selected from those defined in
CC the specification. The primer sets can be used in antisense therapy and
CC in gene therapy. The primers are useful for synthesizing polynucleotides,
CC particularly full-length cDNAs. The primers are also useful for the
CC detection and/or diagnosis of the abnormality of the proteins encoded by
CC the full-length cDNAs. The primers allow obtaining of the full-length
CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
CC AAH13633 to AAH18742 represent human cDNA sequences; AAH92446 to
CC AAH95893 represent human amino acid sequences; and AAH13629 to AAH13632
CC represent oligonucleotides, all of which are used in the exemplification

CC of the present invention.

XX Sequence 1802 BP; 644 A; 298 C; 395 G; 465 T; 0 other;

Query Match 35.4%; Score 1210.2; DB 22; Length 1802;
Best Local Similarity 92.3%; Pred. No. 1.2e-271;
Matches 1287; Conservative 0; Mismatches 103; Indels 5; Gaps 1;

QY 27 gatctcagagtgagcatcatcgattggttatgct-----ttaccattttattagaagact 81
DB 408 gactcgcagagagcatcatcgattggttatgctattgcttaccattttattagaagatt 467
QY 82 atgaatggcagcaaaatttttagaagagtttagaagaaacacagacagacatctcttgata 141
DB 468 atgaatggcagcaaaagatttttagaagaatttttagaagaaacacagacagacatcccttgaca 527
QY 142 agtggattatgaatatagtagaactctcttatatcagaatcaagtctcttcgggaagcag 201
DB 528 agtggattatgaatatagtagaactctcttatatcagaatcaagtctcttcgggaagcag 587
QY 202 gtctttatagaagccctggacatctttgtacatatgaaagcagattttgtataaac 261
DB 588 gtctctatagaagccttggacatctttgtacatatgaaagcagattttgtataaac 647
QY 262 ttgctgttgaagaacccaaaggggaactctctgttcagttgtgtctgtttggaagatgctg 321
DB 648 ttgctgttgaagaacccaaaggggaactctctgttcagttgtgtctgtttggaagatgctg 707
QY 322 ctgacgttttatagaggttacagagaggaatcctgaaattgggcctattacaaaggct 381
DB 708 cagatgttttatagaggttgcaagagagaaatcctgaaactgggcctattacaaaggct 767
QY 382 tagaanaagcactgaagcagcctaataatgttagaagcgttaaaaatatataggaagcct 441
DB 768 tgganaagcactgaagcagcctaataatgttagaagcgttaaaaatatataggaagcct 827
QY 442 ggaactaaatccccggggaactctgccaagaagactcccttaaaacttttttctctgag 501
DB 828 agactaaatatccccggggaactctgccaagaagactcccttaaaacttttttctctgag 587
QY 502 agaaqtttaagagagttgttgataggttctcctaagatgaatttcagaaggggttgcacac 561
DB 898 agaaqtttaagagagttgttgataggttctcctaagatgaatttcagaaggggttgcacac 947
QY 562 ctgtcttcaatcccttgaggtcttttatacagagataagagagaggttgcaatctagaaag 621
DB 948 cagttctcaatcccttgaggtcttttatacagagataagagagaggttgcaatctagaaag 1007
QY 622 aactagtagttggttatgaaactctcttaaaaagttgtgcctatttaacccccaatgatg 681
DB 1008 agttagtagtaggttatgaaactctcttaaaaagttgtgcctatttaacccccaatgatg 1067
QY 682 atggaagaggaggaacccccaacacattacttttggttccagactattttggcagacatt 741
DB 1068 atggaagaggaggaacccccaacacattacttttggttccagactattttggcagacatt 1127
QY 742 atgataaaattggttcagccatccattgctctggaataacataaaactgcaattgaaagta 801
DB 1128 atgataaaattggttcagccatccattgctctggaataacataaaactgcaattgaaagta 1187
QY 802 caccacattgatagaactcttttttgaagaactaaatctataagcatgctggggaata 861
DB 1188 caccacattgatagaactcttttttgaagaactaaatctataagcatgctggggaata 1247
QY 862 ttaagaagctgcccaggttgatgaggaagccagccctggacacagacagacagattta 921
DB 1248 ttaagaagctgcaaggttgatgaggaagccagccctggacacagacagacagattta 1307
QY 922 ttaattccaagttgcaaaatacatgtttaaagccacacctgattaaagaggtctgaagaa 981
DB 1308 tcaactccaaattgtgcaaaatacatgtttaaagccacacctgattaaagaggtctgaagaa 1367

QY 982 tgtgttccaagtattacagagggaagaaacttcagcggttagagaaactgaatgaatgcagt 1041
DB 1368 tgtgttccaagtattacagagggaagaaacttcagcggttagagaaactgaatgaatgcagt 1427
QY 1042 gatatgtgttccagacagagtgtgctcaggcaatacaagcaatgaacaaattttggtgaag 1101
DB 1428 gatatgtgttccaaacagaaatgtgccaggcttataaagcaatgaataaattttggtgaag 1487
QY 1102 cacttcaaaatgtcatgaaattgaagagacatttttatagaatacctgactgaccagtttg 1161
DB 1488 cacttcaaaatgtcatgaaattgaagagacatttttatagaatacctgactgaccagtttg 1547
QY 1162 actttcatcatactgtatgaggaagatcacctcttagatcatatgttgacttattataaac 1221
DB 1548 actttcatcatactgtatgaggaagattacccttagatcatatgttgacttattataaac 1607
QY 1222 tagaagatgtacttcagacagacatcccttttcttcaagcagcgagaatttctattgaga 1281
DB 1608 tagaagatgtacttcagacagacatcccttttcttcaagcagcgagaatttctattgaga 1667
QY 1282 tctatttgaagcttcatgacaacccctctgacagatgagaaacacgagcaggaattgctatgata 1341
DB 1668 tctatttgaagcttcatgacaacccctctgacagatgagaaacacgagcaggaattgctatgata 1727
QY 1342 cagcaaacatgtctgacaaagagctaaagaactgcgttaatacaaaagagagctcaaa 1401
DB 1728 cagcaaacatgtctgacaaagagctaaagaactgcgttaatacaaaagagagctcaaa 1787
QY 1402 agaaagcccagattg 1416
DB 1788 agaaagcccagattg 1802

RESULT 7
AAH14477
ID AAH14477 standard; cDNA; 2859 BP.
XX
AC AAH14477;
XX
DT 26-JUN-2001 (first entry)
XX
DE Human cDNA sequence SEQ ID NO:11977.
XX
KW Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.
XX
OS Homo sapiens.
XX
PN EP1074617-A2.
XX
PD 07-FEB-2001.
XX
PF 28-JUL-2000; 2000EP-0116126.
XX
PR 29-JUL-1999; 99JP-0248036.
PR 27-AUG-1999; 99JP-0300253.
PR 11-JAN-2000; 2000JP-0118776.
PR 02-MAY-2000; 2000JP-0183767.
PR 09-JUN-2000; 2000JP-0241899.
XX
PA (HELI-) HELIX RES INST.
XX
PI Ota T, Isogai T, Nishikawa T, Hayashi K, Salto K, Yamamoto J;
PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX
XX WPI; 2001-318749/34.
XX
PT Primer sets for synthesizing polynucleotides, particularly the 5602
PT full-length cDNAs defined in the specification, and for the detection
PT and/or diagnosis of the abnormality of the proteins encoded by the
XX full-length cDNAs -
PS Claim 8; SEQ ID 11977; 2537pp + CD ROM; English.
XX

The present invention describes primer sets for synthesising 5602 full-length cDNAs defined in the specification. Where a primer set comprises: (a) an oligo-dT primer and an oligonucleotide complementary to the complementary strand of a polynucleotide which comprises one of the 5602 nucleotide sequences defined in the specification, where the oligonucleotide comprises at least 15 nucleotides; or (b) a combination of an oligonucleotide comprising a sequence complementary to the complementary strand of a polynucleotide which comprises a 5'-end sequence and an oligonucleotide comprising a sequence complementary to a polynucleotide which comprises a 3'-end sequence, where the oligonucleotide comprises at least 15 nucleotides and the combination of the 5'-end sequence/3'-end sequence is selected from those defined in the specification. The primer sets can be used in antisense therapy and in gene therapy. The primers are useful for synthesising polynucleotides particularly full-length cDNAs. The primers are also useful for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs. The primers allow obtaining of the full-length cDNAs easily without any specialised methods. AAH03166 to AAH13628 and AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632 represent oligonucleotides, all of which are used in the exemplification of the present invention.

XX sequence 2859 BP: 921 A; 476 C; 547 G; 915 T; 0 other;
SO

Query Match	32.9%	Score 1126;	DB 22;	Length 2859;
Best Local Similarity	90.3%	Pred. No. 5.6e-252;		
Matches 1407: Conservative	0;	Mismatches 105;	Indels 48;	Gaps 17;

[illegible]

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T. J. J. J. J.

RESULTS	0
AAH16424	
ID	AAH16424 standard; cDNA; 1985 BP.
XX	
XX	
AC	AAH16424;
XX	
DT	26-JUN-2001 (first entry)
XX	
DE	Human cDNA sequence SEQ ID NO:15407.
XX	
XX	
XX	Humao. primer. detection; diagnosis; antisense therapy; gene therapy; ss

RESULT 9
AAH81664/c
ID AAH81664 standard; DNA; 710 BP.
XX AC
XX AAH81664;
XX 21-SEP-2001 (first entry)
XX Human differential transcription-associated cDNA SEQ ID 173.
XX DE
XX Differential transcription; human; rat; tumour cell; cytostatic;
KW Ras modulator; Class II tumour suppressor gene; gene therapy; ss.
XX OS
XX Homo sapiens.
XX WO200157058-A2.
XX 09-AUG-2001.
XX 31-JAN-2001; 2001WO-EP01003.
XX 31-JAN-2000; 2000DE-1004102.
XX (META-) METAGEN GES GENOMFORSCHUNG MBH.
XX Rosenthal A, Hinzmann B, Schaefer R, Zuber J, Tchernitsa O;
PI Grips M, Hellriegel M, Schmitz A, Sers C;
XX WPI; 2001-483415/52.
XX Nucleic acids differentially expressed between tumor and normal cells,
PT useful for diagnosis or therapy of tumors and for screening active
PT agents
XX Disclosure; Page 376; 579pp; German.
XX This invention describes a nucleic acid (I) with differential expression
CC between tumor and normal cells and which has cytostatic activity. (I)
CC work as modulators of Ras activity by inducing expression of tumour
CC suppressor genes. (I), and polypeptides encoded by them, are useful as
CC targets for diagnosis or therapy and in screening to determine the
CC effects of an active compound (potential pharmaceutical) on a cell line,
CC particularly for diagnosis and treatment of tumors, especially by
CC modulating expression of (I) (by gene therapy, antisense RNA or ribozyme
CC methods) or by modulating the amount and/or location of (I)-encoded
CC polypeptides (by administration of the polypeptide or its activator,
CC antibody (optionally as a conjugate) or inhibitor). The method allows
CC identification of many Class II tumour suppressor genes (i.e. genes that
CC are not primary targets for tumour-initiating mutations).
CC AAH81492-AAH82376 represent the human and rat derived nucleic acid
CC fragments described in the method of the invention.
XX SQ
Sequence 710 BP; 170 A; 146 C; 121 G; 267 T; 6 other;

Query Match 18.1%; Score 617.8; DB 22; Length 710;
Best Local Similarity 92.4%; Pred. No. 5.8e-134;
Matches 657; Conservative 0; Mismatches 53; Indels 1; Gaps 1;

Oy 714 tgggtccagtactattggccacagcattatgatataaaattggtcagccatccattgctctg 773
Db 710 TGGGTCAAGTACTACTTGGCCACCACTATGACAAAATTGGTCAGCCACTTANTGCTNTG 651

Oy 774 gaatacataataactgaattgaagtagacacccaacttagaactcttcttctgtataaa 833
Db 650 GAGTACATAAATACTGCTATTGAGNAGTACA-CTACATTAATAGACNTCTTCTCGTGANA 592

Oy 834 gctaaatctataagcagctgggaatattataaagaagctgcaggtggatggatgaagcc 893
Db 591 GCTAAATCTATAGCACTGCTGGAATATTAAAGAAGCTGCAAGGTGGATGGATGAGGCN 532

894 caggccctggacacagcagacagattatttaattccaagtgtgcaaaatacatgttataaa 953
531 CAGGCTTGGACACAGCAGACAGATTATCAACTCCAAATGTCAAAATACATGCTAAAA 472
954 gccaacctgattaaagagcgtgaagaattgttccaagtttacgaggggaagaaactca 1013
471 GCCAACCTGATTAAAGAAAGCTGAAGAAATGTCTCAAAAGTTTACAGGGGAAGAACATCA 412
1014 gcggtagaagaacctgaatgaatgcagtgatgtggttccagacagagtgctcaggcca 1073
411 GCGGTAGAGAAATTTGAATGAATGCAGTCGATGTGGTTCCAAACAGAAATGTGCCAGGCT 352
1074 tacaagaacaatgaacaaatttgggtgaagcacttaagaataatgcataaattgagagacat 1133
351 TATAAGCAATGAATAAATTTGGTGAAGCAGCTTAAGAAATGTCAATGAGATTGAGAGACAT 292
1134 ttatagaataatcaccgatgaccaggtttgacttccatactactatgaggaagatcacc 1193
291 TTTATAGAAATCACTGATGACCACTTTGACTTTTACATACATCTATGAGGAAGATTACC 232
1194 cttagatcatatgtggacttattaaaactagaagatgacttcacagcagcattccatttac 1253
231 CTAGATCATATGTGGACTTATTAAACTAGAAAGATGTACTTCGACAGCATCCATTTTAC 172
1254 tcaaaagcagcagagaattgtctattgaagatctatttgaagcttccatgacacccctctgaca 1313
171 TTCAGGCAGCAAGAATTGCTATAGAGATCTATTTTGAAGCTTCATGACACCCCTTACA 112
1314 gatgagaacaaagacacagcagctgatacagacaaacatgtctgacaaagagcgttaagaaa 1373
111 GATGAGAAATAAGAACACAGCAAGCTGATACAGCAAAACATGTCTGACAAAGAGCTAAGAAG 52
1374 ctgcgtataataacaaagaagcgtcctcaaaagaagcccgagattgaagaagag 1424
51 CTACGTAATAAACAAGAAGAGCTCAAAAGAAGCCAGATAGAGAAGAAGAG 1

RESULT 10
AAH06489
ID AAH06489 standard; cDNA; 790 BP.
XX AC
XX AAH06489;
XX 26-JUN-2001 (first entry)
XX Human cDNA clone (5'-primer) SEQ ID NO:3324.
XX Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.
XX Homo sapiens.
XX EP1074617-A2.
XX 07-FEB-2001.
XX 28-JUL-2000; 2000EP-0116126.
XX 29-JUL-1999; 99JP-0248036.
XX 27-AUG-1999; 99JP-0300253.
XX 11-JAN-2000; 2000JP-0118776.
XX 02-MAY-2000; 2000JP-0183767.
XX 09-JUN-2000; 2000JP-0241899.
XX (HELI-) HELIX RES INST.
XX Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX WPI; 2001-318749/34.
XX Primer sets for synthesizing polynucleotides, particularly the 5602
PT full-length cDNAs defined in the specification, and for the detection
PT and/or diagnosis of the abnormality of the proteins encoded by the

proteins, 7 transmembrane receptors, ATPases associated with various cellular activities (AAA), eukaryotic aspartyl proteases, GATA family of transcription factors, G-protein alpha subunit, phospholipase C, diacylglycerol binding proteins, protein kinase, protein phosphatase 2C, protein tyrosine phosphatase, trypsin, wnt family of developmental signalling proteins and WW/rsp5/WWP domain containing proteins. The encoded polypeptides also have a functional domain selected from Ank repeat, basic region plus leucine zipper transcription factors, bromodomain, EF-hand, SH3 domain, WD domain/G-beta repeats, zinc finger (C2H2 type), zinc finger (CCHC class), and zinc-binding metalloprotease domain. The polynucleotides encode polypeptides with similarity to known protein families and are predicted to have similar properties. The novel polynucleotides can be used to develop products for use as therapeutic agents and in forensics, genetic analysis, mapping and diagnostic applications. In particular, the product can be used for the detection and management of cancers. They can be used for treating e.g. cervical cancers, melanomas, colorectal adenocarcinomas, Wilms' tumour, sarcomas, retinoblastoma, myosarcomas, lung carcinomas, leukemias, such as chronic myelogenous leukemia, promyelocytic leukemia, monocytic leukemia, and myeloid leukemia, and lymphomas such as histiocytic lymphoma, anhydric hereditary ectodermal dysplasia, congenital alveolar dysplasia, epithelial dysplasia of the cervix, fibrous dysplasia of bone, and mammary dysplasia, hyperplasias, e.g. endometrial, adrenal, breast, prostate or thyroid hyperplasias or pseudoepitheliomatous hyperplasia of the skin.

Sequence 1225 BP; 400 A; 203 C; 240 G; 379 T; 3 other;

Query Match 16.5%; Score 565; DB 20; Length 1225;
Best Local Similarity 89.7%; Pred. NO. 1.3e-121;
Matches 749; Conservative 1; Mismatches 51; Indels 34; Gaps 12;

QY 2604 ggatttttccactgagcacaagaagttgtgggttttagctatcctgattctgtac 2663
DB 1 ggatttttccactgagcacaagaagttgtgggttttagctatcctgattctgtac 60
QY 2664 ggggtgtgtgatgaccatgaggaatgcaatgtgaatcactgtgtacagagccgtcta 2723
DB 61 ggggtgtgtgat-tctgaccatgaggaatgcaatgtgaatcactgtgtacagagaaacctta 119
QY 2724 caacacatgtctgacgtgttagagactgggacacatagctaccagc-ggattaaagttaa 2782
DB 120 caacagatgtctgattgttagaactgggacacatagctaccagc-ggattaaagttaa 179
QY 2783 accatgaggtgttcagtagctgtgtgtgtttccaaaatcactgtacatgatcagttt 2842
DB 180 accatgaggtgttcagtagc-ctgtgtgtttccaaaatcactgtacatgatcagttt 236
QY 2843 ggtgtctgtaccacagtttttaaccgaaggaaccaggtgtggaacatctcaa-tttaac 2901
DB 237 ggtgtctgtaccacagtttttaaccgaaggaaccaggtgtggaacatctcaa-tttaac 296
QY 2902 taaactgtgaagactaaataaacaatgcaacactttatcattgtttggcacaactgt 2961
DB 297 taaactgtgaagactaaataaacaatgcaacactttcagcattgtttggcacaactgt 356
QY 2962 taaactgtgaagactaaataaacaatgcaacactttcagcattgtttggcacaactgt 3021
DB 357 taaactgtgaagactaaataaacaatgcaacactttcagcattgtttggcacaactgt 415
QY 3022 tgaatttttccactgagagtgtaaaagaaactacacatggtgtgaagtacagagc 3081
DB 416 tgaatttttccactgagagtgtaaaagaaactacacatggtgtgaagtacagagc 474
QY 3082 agaactcctgactacattctctgactgtgactgtgagagactaata-tctaaaaacctcagcag 3140
DB 475 agaactcctgactacattctctgactgtgactgtgagagactaata-tctaaaaacctcagcag 534
QY 3141 gcttgtttcacgatatgcag---aaaaagctgtcagtttagatcacctctgggaactttt 3197
DB 535 gcttgtttcacgatatgcagaaaaaaagctgtcagtttagatcacctct-ggaattttt 593

QY 3198 ccacaggtcacaggtttgttaacttaactgaagcccttcatttctaagaataatttctgc 3257
DB 594 ccacaggtcacaggtttgttaacttaactgaagcccttcatttctaagaataatttctgc 653
QY 3258 tcagttgtttcagcagcagcccaagactttgttaatttttaaggcccaagatattttttt 3317
DB 654 tcagttgtttcagcagcagcccaagactttgttaatttttaaggcccaagatattttttt 713
QY 3318 -----caataacagacccagcttttttctcagttacaaatgtaattct 3364
DB 714 ttttttttttttcaataaagacccagcttttttctcagttacaaatgtaattct- 772
QY 3365 ttttttttttttcaataaagacccagcttttttctcagttacaaatgtaattct 3416
DB 773 -----ctttttgttcaataaagacccagcttttttctcagttacaaatgtaattct 823

RESULT 12

AAZ15705 ID AAZ15705 standard; cDNA; 781 BP.
XX AC AAZ15705;
XX 12-OCT-1999 (first entry)
XX Human gene expression product cDNA sequence SEQ ID NO:3174.
XX Human; gene; gene expression product; diagnosis; therapy; probe;
KW detection; mapping; tissue typing; profiling; forensic; cancer;
KW genetic analysis; colorectal cancer; breast cancer; lung cancer; ss.
OS Homo sapiens.
XX WO9398972-A2.
XX 05-AUG-1999.
XX 28-JAN-1999; 99WO-US01619.
XX 03-APR-1998; 98US-0080666.
XX 28-JAN-1998; 98US-0072910.
XX 24-FEB-1998; 98US-0075954.
XX 31-MAR-1998; 98US-0080114.
XX 03-APR-1998; 98US-0080515.
XX (CHIR) CHIRON CORP.
XX (HYSE-) HYSEQ INC.
XX Crkvenjakov R, Dickson M, Drmanac R, Drmanac S;
PI Escobedo J, Garcia PD, Garcia V, Giese K, Innis MA;
PI Jones WL, Kassam A, Kennedy GC, Kita D, Labat I;
PI Lamson G, Leshkowitz D, Pot D, Randazzo F, Reinhard C;
PI Stache-Crain B, Sudduth-Klinger J, Williams LT;
XX WPI; 1999-494092/41.
XX Novel human genes and their expression products which are
XX differentially expressed in different cell types
XX Claim 1; Page 1524-1525; 2479pp; English.

The present invention describes a library of human polynucleotides comprising the sequences given in AAZ12532 to AAZ17779. Also described is a method of detecting differentially expressed genes correlated with the cancerous state of a mammalian cell, comprising detecting at least one differentially expressed gene product in a test sample from a cell suspected of being cancerous, where the gene product is encoded by one of the 5248 polynucleotide sequences given in AAZ12532 to AAZ17779. The polynucleotides can be used as a source of primers and probes, which can be used for a variety of purpose, e.g. detection of expression levels, mapping, tissue typing or profiling, forensics, genetic analysis and detection of polymorphisms. Polypeptides encoded by the polynucleotides can be used for raising antibodies for experimental, diagnostic and

CC therapeutic purposes. The polynucleotides may also be used to construct
CC arrays for diagnostics (which may be used to determine function of an
CC encoded protein); and to detect differences in expression levels between
CC two cells (e.g. to identify abnormal or diseased tissue in a human, to
CC identify a genetic predisposition or susceptibility to a disease such as
CC cancer). The polynucleotides of the invention are especially used in the
CC diagnosis, prognosis and management of colorectal cancer, breast cancer,
CC and lung cancer. The polynucleotides can also be used to screen for
CC peptide analogues and antagonists.
XX
SQ Sequence 781 BP; 263 A; 140 C; 156 G; 205 T; 17 other;

	Query Match	16.3%;	Score 535.6;	DB 20;	Length 781;
	Best Local Similarity	90.8%;	Pred. No. 1.8e-119;		
	Matches 621;	Conservative 0;	Mismatches 60;	Indels 3;	Gaps 3;
QY	597 aaagagaaggtggaacatctagaagaactagtagtggttgataaaactctctataaaagt	656			
Db	94 aagaaaaggtggaacatctagaagaactagtagtggttgataaaactctctataaaagt	153			
QY	657 tgcgcctatttaaccccaatgaatgaagaagggaacacctcccaaccacatactttgg	716			
Db	154 tgcgcctatttaaccccaatgaatgaagaagggaacacctcccaaccacatactttgg	213			
QY	717 gtccagtagctatttgccacagcatctattgataaaattggtcagccatccattgctctgaa	776			
Db	214 gtccnntnctactggtgacacacatctatgacaaaattggtcagccatccattgctctgaa	273			
QY	777 tacataaactgcattgaaagtagtaacacacacattgataagaactctttctgtaaaagt	836			
Db	274 tacataaactgcattgaaagtagtaacacacacattgataagaactctttctgtaaaagt	333			
QY	837 aaaaactataagcatgctgggaatattaaagaagtgccagggtgatggatgaagccag	896			
Db	334 aaaaactataagcatgctgggaatattaaagaagtgccagggtgatggatgaagccag	393			
QY	897 gccctggacacagcagacagatatttatttccaaagtgtgcaaaatgatactgttaaaagcc	956			
Db	394 gccctggacacagcagacagatatttatttccaaagtgtgcaaaatgatactgttaaaagcc	453			
QY	957 aacctgattaaagcgtgaagaaatgtgttccaaagtttacagaggaaggaacttcagcg	1016			
Db	454 aacctgattaaagcgtgaagaaatgtgttccaaagtttacagaggaaggaacttcagcg	513			
QY	1017 gttagagaacctgaaatgaatgcagtgatgtggttccagacagagatggtcaggcagcatac	1076			
Db	514 gttagagaacctgaaatgaatgcagtgatgtggttccagacagagatggtcaggcagcatac	573			
QY	1077 aaagcaatgaacaaatttggtagagcacttaagaatgtcatgaattgagagacatttt	1136			
Db	574 aaagcaatgaacaaatttggtagagcacttaagaatgtcatgaattgagagacatttt	633			
QY	1137 ata-gaaatcaccgatgaccagtttgactttacatactatgtatg-aggaagatcaccc	1194			
Db	634 atagaaatactatgaccagtttgactttacatactatgtatg-aggaagatcaccc	693			
QY	1195 ttatgatcatgttgactttataaacta-gaagatgtacttcagacagatccattttac	1253			
Db	694 ttatgatcatgttgactttataaacta-gaagatgtacttcagacagatccattttac	753			
QY	1254 ttcaagcagcagaattgctatt 1277				
Db	754 ttcaagcagcagaattgctatt 777				

RESULT 13

AAX99053

ID AAX99053 standard; cDNA; 781 BP.

XX AAX99053;

XX AAX99053;

DT 24-SEP-1999 (first entry)

XX

DE

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KW

KW

KW

KW

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KW

KW

KW

XX

OS

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PN

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PD

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Human validated cancer cell derived cDNA #375.

Cancer; human; colon; breast; lung; transmembrane receptor; ATPase;
integral membrane protein; aspartyl protease; GATA family; wnt family;
transcription factor; G-protein alpha subunit; protein phosphatase;
phorbol ester binding protein; diacylglycerol binding protein; trypsin;
protein kinase; tyrosine phosphatase; developmental signalling protein;
WW/rsp5/WWP domain; therapy; forensic; genetic mapping; diagnostic;
detection; treatment; cervical; melanoma; colorectal adenocarcinoma;
Wilm's tumour; retinoblastoma; sarcoma; myosarcoma; lung carcinoma;
leukemia; lymphoma; dysplasia; hyperplasia; endometrium; adrenal;
prostate; ss.

Homo sapiens.

WO9933982-A2.

08-JUL-1999.

22-DEC-1998; 98WO-US27610.

21-DEC-1998; 98US-0217471.

23-DEC-1997; 97US-0068755.

03-APR-1998; 98US-0080664.

21-OCT-1998; 98US-0105234.

27-OCT-1998; 98US-0105877.

(CHIR) CHIRON CORP.

(HYSE) HYSEQ INC.

Crkvenjakov R, Dickson M, Drmanac R, Drmanac S;

Escobedo J, Garcia PD, Garcia V, Giese K, Innis MA;

Jones LW, Kassam A, Kennedy GC, Kita D, Labat I;

Lanson G, Leshkowitz D, Pot D, Randazzo F, Reinhard C;

Stache-Crain B, Sudduth-Klinger J, Williams LT;

WPI; 1999-430243/36.

New isolated human polynucleotides

Claim 1; Page 564; 591pp; English.

This invention describes novel isolated human polynucleotides obtained by screening for differential expression in colon cancer, breast cancer and lung cancer cell lines. The polynucleotides of the invention are represented in AAX98275-X99118 and encode polypeptides of protein families selected from 4 transmembrane segments integral membrane proteins, 7 transmembrane receptors, ATPases associated with various cellular activities (AAA), eukaryotic aspartyl proteases, GATA family of transcription factors, G-protein alpha subunit, phospholipases or diacylglycerol binding proteins, protein kinase, protein phosphatase 2C, signalling proteins and WW/rsp5/WWP domain containing proteins. The encoded polypeptides also have a functional domain selected from Ank repeat, basic region plus leucine zipper transcription factors, bromodomain, EF-hand, SH3 domain, WD domain/G-beta repeats, zinc finger (C2H2 type), zinc finger (CCHC class), and zinc-binding metalloprotease domain. The polynucleotides encode polypeptides with similarity to known protein families and are predicted to have similar properties. The novel polynucleotides can be used to develop products for use as therapeutic agents and in forensics, genetic analysis, mapping and diagnostic applications. In particular, the product can be used for the detection and management of cancers. They can be used for treating e.g. cervical cancers, melanomas, colorectal adenocarcinomas, Wilm's tumour, sarcomas, retinoblastoma, myosarcomas, lung carcinomas, leukemias, such as chronic myelogenous leukemia, promyelocytic leukemia, monocytic leukemia, and myeloid leukemia, and lymphomas such as histiocytic lymphoma, anhydric hereditary ectodermal dysplasia, congenital alveolar dysplasia, epithelial dysplasia of the cervix, fibrous dysplasia of bone, and mammary dysplasia, hyperplasias, e.g. endometrial, adrenal, breast, prostate or thyroid hyperplasias or pseudopitheliomatous hyperplasia of the skin.

[illegible]

Mon Jul 22 09:40:57 2002

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Db 485 tcagcggttagagaatttgaaatgcagtgcatgtgttccaaacagaatgtgcccag 544
QY 1071 gcatacaaaagcaatgaa-caaatttgggtgaagcacttaagaaatgtca-tgaaattgaga 1128
Db 545 gcttataaagcaatgaattaaatttgggtgaagcacttaagaaatgtcattgagattgaga 604
QY 1129 gacattttatagaaatcacccgatga-ccagtttgactttcatatactactgtatga-ggaa 1186
Db 605 gacttttataggaatcactgatgaccagtttgactttcatatactactgtatganggaa 664
QY 1187 gatcaccccttag 1198
Db 665 nattaacccttag 676
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Search completed: July 20, 2002, 02:41:40
Job time: 7818 sec

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ORIGIN		
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	Ratio: 4.778 Gaps: 0	
Percent Similarity:	100.000 Percent Identity: 88.889	
alignment_block:		
US-09-836-410A-1 x AX159092/rev ..		
Align seg 1/1 to reverse of: AX159092 from: 1 to: 50		
17 ArgGlyLeuValProArgLysLeuPro 25		
43 AGAGGGCTGGTCCACGTCAGTGCCG 17		
seq_name: gb_pat:AR126194		
seq_documentation_block:		
LOCUS AR126194	45 bp DNA	linear PAT 16-MAY-2001
DEFINITION Sequence 212 from patent US 6180084.		
ACCESSION AR126194		
VERSION AR126194.1 GI:14112787		
KEYWORDS	Unknown.	
SOURCE	Unknown.	
ORGANISM	Unclassified.	
REFERENCE 1 (bases 1 to 45)		
AUTHORS Ruoslahti, E. and Pasqualini, R.		
TITLE NGR receptor and methods of identifying tumor homing molecules that		
JOURNAL home to angiogenic vasculature using same		
FEATURES Patent: US 6180084-A 212 30-JAN-2001;		
Location/Qualifiers		
1..45		
Source		
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ORIGIN		
alignment_scores:	Quality: 42.00 Length: 14	
	Ratio: 3.818 Gaps: 0	
Percent Similarity:	78.571 Percent Identity: 50.000	
alignment_block:		
US-09-836-410A-1 x AR126194 ..		
Align seg 1/1 to: AR126194 from: 1 to: 45		
91 AspAspGlyLysGluGluProThrThrLeuLeuTrpVal 104		
2 GAAGATCTAGAGGAACCCGCCCTGTGACGGTCAGCTGGGTC 43		
seq_name: gb_pat:A70028		
seq_documentation_block:		
LOCUS A70028	42 bp DNA	linear PAT 07-MAY-1999
DEFINITION Sequence 11 from Patent EP0832972.		
ACCESSION A70028		
VERSION A70028.1 GI:4774476		
KEYWORDS	unidentified.	
SOURCE	unidentified	
ORGANISM	unclassified.	
REFERENCE 1 (bases 1 to 42)		
AUTHORS Silke, N., Lerch, K. and Muheim, A.		
TITLE Cloning, expression and production of tasty peptides		
JOURNAL Patent: EP 0832972-A 11 01-APR-1998;		
Location/Qualifiers		
1..50		
Source		
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    /db_xref="taxon:32644"
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  ORIGIN

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  Ratio: 3.375        Gaps: 1
  Percent Similarity: 80.000      Percent Identity: 53.333

alignment_block:
  US-09-836-410A-1 x A70028 ..
  Align seg 1/1 to: A70028 from: 1 to: 42

243 PheileGluIleThrAspGlnPheHisThrTyrCys 257
1 TTTCTTAGAA...ACCGATTCTCTCTCGACTTCGATTCGCCACTGC 42
seq_name: gb_pat:AX221610

seq_documentation_block:
  LOCUS      AX221610      48 bp      mRNA      linear      PAT 07-SEP-2001
  DEFINITION      Sequence 7052 from Patent WO0159103.
  ACCESSION      AX221610
  VERSION      AX221610.1      GI:15549334
  KEYWORDS
  SOURCE
  ORGANISM
  synthentic construct.
  artificial sequence.
  REFERENCE
  1 (bases 1 to 48)
  AUTHORS      Blatt,L., McSwiggen,J. and Chowrira,B.M.
  TITLE      Method and reagent for the modulation and diagnosis of cd20 and
  JOURNAL      nogo gene expression
  PATENT: WO 0159103-A 7052 16-AUG-2001;
  RIBOZYME PHARMACEUTICALS, INC. (US); Blatt, Lawrence (US);
  McSwiggen, James (US); Chowrira, Bharat M. (US)
  FEATURES
  source      Location/Qualifiers
  1..48
    /organism="synthetic construct"
    /db_xref="taxon:32630"
    /note="Nucleic Acid"
  BASE COUNT      15 a 11 c 12 g 10 t
  ORIGIN

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  Ratio: 2.893        Gaps: 1
  Percent Similarity: 73.684      Percent Identity: 47.368

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381 ProLeuGluAlaIleLysPheLeuThrProLeuLysAsnLeuVally 397
4 CCCTTGAGGAA.....ACTCCCTCAAGGACATCGTCG 38
397 sAsnLys 399
39 GGATAAA 45
seq_name: gb_pr:HUMRPY59

seq_documentation_block:
  LOCUS      HUMRPY59      48 bp      mRNA      linear      PRI 27-SEP-2001
  DEFINITION      Homo sapiens mRNA for ribosomal protein L24, partial cds.
  ACCESSION      D28400
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VERSION      D28400.1      GI:461272
KEYWORDS
SOURCE      Homo sapiens lymphoma cell_line:U937 cdNA to mRNA,
clone_lib:U937/pkAl clone:HP00302.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 48)
AUTHORS      Kato,S., Sekine,S., Oh,S.W., Kim,N.S., Umezawa,Y., Abe,N.,
Yokoyama-Kobayashi,M. and Aoki,T.
TITLE      Construction of a human full-length cdNA bank
JOURNAL      Gens. 150 (2), 243-250 (1994)
MEDLINE
95121910
REFERENCE
2 (bases 1 to 48)
AUTHORS      Kato,S.
TITLE      Direct Submission
JOURNAL      Submitted (03-FEB-1994) Seishi Kato, Research Institute of National
Rehabilitation Center for the Disabled, Department of
Rehabilitation Engineering; 4-1 Namiki, Tokorozawa, Saitama
359-8555, Japan (E-mail:seishi@rehab.go.jp,
Tel:81-42-995-3100(ex.2568), Fax:81-42-995-3132)
FEATURES
  source      Location/Qualifiers
  1..48
    /organism="Homo sapiens"
    /db_xref="taxon:9606"
    /clone="HP00302"
    /cell_line="U937"
    /tissue_type="lymphoma"
    /clone_lib="U937/pkAl"
  1..42
    /replace="ttttcttttc"
  1..10
    /replace="cttttttttc"
  1..10
    /replace="cttttttttc"
  1..10
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  1..10
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  1..10
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    /product="ribosomal protein L24"
    /protein_id="BAA05766.1"
    /db_xref="GI:4433237"
    /translation="MK"
  BASE COUNT      5 a 13 c 10 g 20 t
  ORIGIN

alignment_scores:
  Quality: 40.00      Length: 15
  Ratio: 2.857        Gaps: 0
  Percent Similarity: 93.333      Percent Identity: 46.667

alignment_block:
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  Align seg 1/1 to reverse of: HUMRPY59 from: 1 to: 48

578 ValAsnGlyAspSerAlaGluThrGluGluLeuAlaAsnGlu 592
48 CITCATGGCGGACAGCTCCACGGAACACAAAGATGGCGGAAGAA 4
seq_name: gb_pat:AX147194

seq_documentation_block:
  LOCUS      AX147194      50 bp      DNA      linear      PAT 08-JUN-2001
  DEFINITION      Sequence 32 from Patent WO0136682.
  ACCESSION      AX147194
  VERSION      AX147194.1      GI:14346365
  KEYWORDS
  SOURCE      synthetic construct.
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36 CCGG 39

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ORGANISM synthetic construct
artificial sequence.
REFERENCE 1 (bases 1 to 50)
AUTHORS Chenchik,A., Munishkin,A. and Simonenko,P.
TITLE Long oligonucleotide arrays
JOURNAL Patent: WO 0136682-A 32 25-MAY-2001;
Clontech Laboratories Inc. (US)
FEATURES
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        /organism="synthetic construct"
        /db_xref="taxon:32630"
        /note="synthetic oligonucleotide"
BASE COUNT 17 a 6 c 16 g 11 t
ORIGIN
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    Percent Similarity: 100.000 Percent Identity: 66.667
alignment_block:
    US-09-836-410A-1 x AX147194/rev ..
Align seg 1/1 to reverse of: AX147194 from: 1 to: 50
428 AlaileAspSerHisProTrrpLeu 436
|||||:|||||:|||||:|||||
44 GCACCTTTCTCAAGTCACCCCTTGCGTG 18
seq_name: gb_pat:AX222245
seq_documentation_block:
    LOCUS AX222245 48 bp mRNA linear PAT 07-SEP-2001
    DEFINITION Sequence 7687 from Patent WO0159103.
    ACCESSION AX222245
    VERSION AX222245.1 GI:15549969
    KEYWORDS
        synthetic construct.
        synthetic construct.
        artificial sequence.
    ORGANISM
        synthetic construct.
    REFERENCE
        1 (bases 1 to 48)
        Blatt,L., McSwiggen,J. and Chowrira,B.M.
        Method and reagent for the modulation and diagnosis of cd20 and
        nogo gene expression
    JOURNAL Patent: WO 0159103-A 7687 16-AUG-2001;
        RIBOZYME PHARMACEUTICALS, INC. (US) ; Blatt, Lawrence (US) ;
        McSwiggen, James (US) ; Chowrira, Bharat M. (US)
    FEATURES
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            1..48
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            /note="Nucleic Acid"
BASE COUNT 12 a 16 c 11 g 9 t
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alignment_scores:
    Quality: 39.50 Length: 18
    Ratio: 3.038 Gaps: 1
    Percent Similarity: 72.222 Percent Identity: 50.000
alignment_block:
    US-09-836-410A-1 x AX222245 ..
Align seg 1/1 to: AX222245 from: 1 to: 48
380 ThrProLeuGluAlaLeuLysPheLeuThrProLeuLysAsnLeuVa 396
|||||:|||||:|||||:|||||:|||||:|||||
1 ACACCACTGGAGGAA.....ACTCCCTTCAAGGACATCGT 35
396 lLys 397
|:::

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seq_name: gb_pat:AX111996
seq_documentation_block:
    LOCUS AX111996 30 bp DNA linear PAT 01-MAY-2001
    DEFINITION Sequence 9 from Patent WO0125439.
    ACCESSION AX111996
    VERSION AX111996.1 GI:13938904
    KEYWORDS
        synthetic construct.
        synthetic construct.
        artificial sequence.
    ORGANISM
        1 (bases 1 to 30)
        Bonello,J.F., Rogowsky,P. and Perez,P.
        Plant seed endosperm-specific promoter
        TITLE Patent: WO 0125439-A 9 12-APR-2001;
        Biogemma (FR)
    FEATURES
        Location/Qualifiers
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                1..30
                /organism="synthetic construct"
                /db_xref="taxon:32630"
                /note="SEQUENCE DESCRIPTION artificielle:oligonucleotide"
BASE COUNT 9 a 6 c 6 g 9 t
ORIGIN
alignment_scores:
    Quality: 39.00 Length: 10
    Ratio: 4.333 Gaps: 0
    Percent Similarity: 90.000 Percent Identity: 70.000
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Align seg 1/1 to: AX111996 from: 1 to: 30
230 GlyGluAlaLeuLysLysCysHisGluIle 239
|||||:|||||:|||||:|||||:|||||
1 GGGGAAGCTTTACATTCTTGCCATAACATA 30
seq_name: gb_pat:AR123920
seq_documentation_block:
    LOCUS AR123920 47 bp DNA linear PAT 16-MAY-2001
    DEFINITION Sequence 4 from patent US 6171823.
    ACCESSION AR123920
    VERSION AR123920.1 GI:14109281
    KEYWORDS
        Unknown.
        Unknown.
    ORGANISM
        Unclassified.
    REFERENCE
        1 (bases 1 to 47)
        Woldike,H.Fabricius, and Hastrup,S.
        Process for producing extracellular proteins in bacteria
        TITLE Patent: US 6171823-A 4 09-JAN-2001;
        JOURNAL Location/Qualifiers
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        source
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BASE COUNT 6 a 18 c 12 g 11 t
ORIGIN
alignment_scores:
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    Ratio: 4.333 Gaps: 0
    Percent Similarity: 100.000 Percent Identity: 66.667
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Align seg 1/1 to reverse of: AR123920 from: 1 to: 47

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355 LysLysAspAspAspGluGluIle 363
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27 GAGAGGACGACGATGATAAGAGGTC 1

seq_name: gb_pat:129948

seq_documentation_block:

LOCUS I29948 50 bp DNA linear PAT 06-FEB-1997

DEFINITION Sequence 12 from patent US 5578478.

ACCESSION I29948

VERSION I29948.1 GI:1820739

KEYWORDS

SOURCE

ORGANISM

REFERENCE

1 (bases 1 to 50)

AUTHORS Rambossek, J., Piddington, C.S., Kovacevich, B.R., Young, K.D. and

Denome, S.A.

TITLE Recombinant DNA encoding a desulfurization biocatalyst

JOURNAL Patent: US 5578478-A 12 26-NOV-1996;

FEATURES

Location/Qualifiers

1..50

/organism="unknown"

BASE COUNT 17 a 13 c 12 g 8 t

ORIGIN

alignment_scores:

Quality: 39.00 Length: 9

Ratio: 4.875 Gaps: 0

Percent Similarity: 88.889 Percent Identity: 66.667

alignment_block:

US-09-836-410A-1 x I29948/rev ..

Align seg 1/1 to reverse of: I29948 from: 1 to: 50

553 AlaSerCysHisLysLeuPheProTyr 561
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43 GCGGCTTGTCATGCTGTTCTCAT 17

seq_name: gb_pat:AX223522

seq_documentation_block:

LOCUS AX223522 48 bp mRNA linear PAT 07-SEP-2001

DEFINITION Sequence 8964 from Patent WO0159103.

ACCESSION AX223522

VERSION AX223522.1 GI:15551246

KEYWORDS

SOURCE

ORGANISM

REFERENCE

1 (bases 1 to 48)

AUTHORS Blatt, L., McSwiggen, J. and Chowrira, B.M.

TITLE Method and reagent for the modulation and diagnosis of cd20 and

nogo gene expression

JOURNAL Patent: WO 0159103-A 8964 16-AUG-2001;

FEATURES

Location/Qualifiers

1..48

/organism="synthetic construct"

/db_xref="taxon:32630"

/note="Nucleic Acid"

BASE COUNT 13 a 13 c 12 g 10 t

ORIGIN

alignment_scores:

Quality: 38.50 Length: 21

Ratio: 2.406 Gaps: 1

Percent Similarity: 76.190 Percent Identity: 38.095

alignment_block:

US-09-836-410A-1 x AX223522 ..

Align seg 1/1 to: AX223522 from: 1 to: 48

380 ThrProLeuGluAlaIleLysPheLeuThrProLeuLysAsnLeuVa 396
:::|||||
1 AGCCCATTTGGAGGAA.....ACTCCCTTCAAGGACATCGT 35

396 lLysAsnLysIle 400
|:::|||||
36 CCGGGATATCTG 48

seq_name: gb_pat:AX229410

seq_documentation_block:

LOCUS AX229410 48 bp mRNA linear PAT 10-SEP-2001

DEFINITION Sequence 2782 from Patent WO0157206.

ACCESSION AX229410

VERSION AX229410.1 GI:15558551

KEYWORDS

SOURCE

ORGANISM

REFERENCE

1 (bases 1 to 48)

AUTHORS Fattaey, A.R., Jarvis, T., Mcswlggen, J., Booher, R.N. and Holman, P.S.

TITLE Method and reagent for the inhibition of checkpoint kinase-1 (chk

1) enzyme

JOURNAL Patent: WO 0157206-A 2782 09-AUG-2001;

FEATURES

Location/Qualifiers

1..48

/organism="synthetic construct"

/db_xref="taxon:32630"

BASE COUNT 11 a 14 c 14 g 9 t

ORIGIN

alignment_scores:

Quality: 38.50 Length: 21

Ratio: 2.567 Gaps: 1

Percent Similarity: 71.429 Percent Identity: 38.095

alignment_block:

US-09-836-410A-1 x AX229410 ..

Align seg 1/1 to: AX229410 from: 1 to: 48

380 ThrProLeuGluAlaIleLysPheLeuThrProLeuLysAsnLeuVa 396
|:::|||||
1 ACCCTCGCGGAGGAA.....ACTCCCTTCAAGGACATCGT 35

396 lLysAsnLysIle 400
|:::|||||
36 CCGGGATGAGTTG 48

us-09-836-410a-1.p2n15to50.rge

Mon Jul 22 09:40:55 2002

OM of: US-09-836-410A-1 to: N_Geneseq_032802.* out_format : pfs
Date: Jul 20, 2002 4:52 AM
About: Results were produced by the GenCore software, version 4.5
Copyright (c) 1993-2000 CompuGen Ltd.

```

Command line parameters:
-MODEL=frame+pn2n.model
-o=/cgn2_1/USPTO_spool/US09836410/runat_l8072002_l15033_29721/app_query.fasta_1.660
-DB=N_Genesec 032802 -QFMT=fastap -SUFFIX=pn2n15to50.rmg
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-LCOPEXT=0.000 -QGAPOP=4.500 -QGAPEXT=0.050 -XGAPOP=10.000
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-YGAPEXT=0.500 -DELOP=6.000 -DELEXT=7.000 -START=1
-MATRIX=blosum62 -TRANS=human40.cdi -LIST=45 -DocalIGN=200
-THR_SCORE=pct -THR_MAX=100 -THR_MIN=0 -ALIGN=15 -MODE=LOCAL
-OUTFMT=pfs -NORM=ext -HEAPSIZE=500 -MINLEN=15 -MAXLEN=50
-USER=US09836410@cgn1_1_188 -NCPU=6 -ICPU=3 -LONGLOG
-DEV_TIMESOUT=120 -MIN_TIMESOUT=30 -NO_XLPXY -WAIT -THREADS=1

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Search information block:

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Search information block:
Query: US-09-836-410A-1
Query length: 593
Database: N_Geneseq_032802.*
Database sequences: 1736436
Database length: 858457221
Search time (sec): 200.300000
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score_list:

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/SDSI/cgdata/hold-geneseq/geneseq-emb1/NA1999.DAT:AAAY05908	+	41.00	84.02	2.5e+04	
/SDSI/cgdata/hold-geneseq/geneseq-emb1/NA1999.DAT:AAH15480	+	41.00	84.02	2.5e+04	
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seq_name: /SIDSL/gcgdata/hold-geneseq/geneseqn-embl/NA2001A.DAT:AAH27115

seq_documentation block:

ID AAH27115 standard; DNA; 49 BP.

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DT 06-AUG-2001 (first entry)

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PCR primer used for FEN endonuclease gene cloning.

KW Cleavage structure; target sequence detection; flap endonuclease;

KW FEN; PCR primer; ss.

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pyrococcus furiosus

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PN WO200132922-A2.

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PD
10-MAY-2007

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PF 27-OCT-2000; 2000WO-US29663.

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SNEE; T00-67
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PA (STRA-) STRATAGENE.

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301ge JA;

DR WPI; 2001-328805/34.

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PS Example 2: Page 55: 81pp: Englishb.

This invention relates to a method for generating a signal indicative of the presence of a target nucleic acid sequence in a sample. The method comprises the formation of a cleavage structure through the incubation of a sample comprising a target nucleic acid sequence and a nucleic acid polymerase and cleaving the cleavage structure with a 5' exonuclease-1 or flap endonuclease (FEN) to generate the signal. The method is used for the detection and quantification of a target nucleic acid sequence. The present sequence represents a PCR primer used to amplify the Pyrococcus furiosus FEN endonuclease gene sequence. The PCR product is used in an example illustrating the method of the invention.

XX
Sequence 49 BP; 18 A; 8 C; 14 G; 9 T; 0 other;

alignment_scores:

Quality:	44.00	Length:	16
Ratio:	3.385	Gaps:	0
Percent Similarity:	81.250	Percent Identity:	50.000

alignment_block:

US-09-836-410A-1 x AAH27115

Align seq 1/1 to: AAH27115 from: 1 to: 49

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359 AspAspGluGluIleGlyGlyProLysGluGluLeuIleProGluLys 374

1 GAGGACCAACATCCCTCCCATTTCTCACCATTATTCACATTTTACAAAT

1 GAGGCGCCGGCCTGGTGTCTCCCAATGGGTGAGATATAACCAAGAA 48

seq_name: /SIDS1/gcgdata/hold-geneseq/geneseqn-emb1/NA2001A.DAT:AAF57073

See document on block.

```

seqname: /stgs1/ccadata/bold-geneseq/geneseq-emb1/NA2000.DAT:AAA08530

```

seq_documentation_block:
 ID AAA08530 standard; DNA; 50 BP.
 AC AAA08530;
 XX
 DT 19-JUL-2000 (first entry)
 XX
 DE Oligonucleotide encoding C-terminal portion of the alpha factor signal
 DE sequence and mutated trypsinogen leader sequence.
 XX
 KW C-terminal alpha factor signal sequence; trypsinogen; leader sequence;
 KW analogue; mutated bovine trypsinogen; recombinant protein production;
 KW inactive zymogen; ss.
 XX
 OS Synthetic.
 XX
 PN WO200017332-A1.
 XX
 PD 30-MAR-2000.
 XX
 PF 15-SEP-1999; 99WO-US21047.
 XX
 PR 21-SEP-1998; 98US-0101213.
 XX
 PA (ELIL) LILLY & CO ELI.
 XX
 PI Hanqueler JM, Hershberger CL, Desplancq D, Larson JL, Rosteck PR;
 XX
 DR WPI: 2000-283565/24.
 XX
 PT New trypsinogen analog useful for the production of recombinant trypsin
 PT has a modified leader sequence not cleavable by trypsin or trypsin-like
 PT enzymes
 XX
 PS Example 1; Page 29; 56pp; English.
 XX
 CC Trypsinogen was fused directly to the C-terminus of the alpha factor
 CC without a Glu-Ala-Glu-Ala linker peptide. Oligonucleotides AAA08529-30
 CC encoding a C-terminal portion of the alpha factor signal sequence and
 CC the Val(Asp)5 leader sequence were synthesized. The wild type bovine
 CC trypsinogen was mutated to destroy the trypsin cleavage site. The lys
 CC residue present in the leader sequence of the native bovine trypsinogen
 CC protein was mutated to an Asp residue. The specification claims an
 CC isolated trypsinogen analogue comprising a protein having trypsin
 CC activity and a leader sequence having at least two amino acids which
 CC are not Lys or Arg. A recombinantly produced trypsin (AA91926) is also
 CC claimed. The trypsin derived from the recombinant trypsinogen is useful
 CC for the characterization of other proteins, and in the manufacture of
 CC other recombinant bioproducts, for example to cleave leader sequences
 CC from small recombinant proteins expressed initially as fusion proteins.
 CC The present method provides for expression of an inactive zymogen form
 CC that is soluble and properly folded yet is not activated until after
 CC purification from fermentation broth or cell extracts. This is
 CC accomplished through the expression of a single chain trypsinogen
 CC analogue where the leader sequence is modified such that it lacks a
 CC trypsin-like enzyme cleavage site. Specifically the trypsinogen
 CC analogues of the present invention lack a lysine or arginine in the
 CC N-terminal leader sequence of the protein to prevent auto-activation or
 CC activation by endogenous host cell enzymes.
 XX
 SQ Sequence 50 BP; 12 A; 17 C; 6 G; 15 T; 0 other;

alignment_scores:
 Quality: 43.00 Length: 12
 Ratio: 4.300 Gaps: 0
 Percent Similarity: 83.333 Percent Identity: 58.333
 alignment_block:
 US-09-836-410A-1 x AAA08530/rev ..
 Align seg 1/1 to reverse of: AAA08530 from: 1 to: 50

354 LysLysLysAspAspAspGluGluIleGlyGly 365
 |||||
 49 AAAAGAGTCGACGATGATGACGATATCGTTGGAGGT 14
 seq_name: /SIDSI/gc9data/hold-geneseq/geneseq-emb1/NA2001A.DAT:AAI75479
 seq_documentation_block:
 ID AAI75479 standard; DNA; 50 BP.
 XX
 AC AAI75479;
 XX
 DT 09-NOV-2001 (first entry)
 XX
 DE Human silent SNP containing nucleic acid SEQ:2420.
 XX
 KW Human; single nucleotide polymorphism; SNP; genome; gene therapy;
 KW protein therapy; vaccine; probe; diagnostic assay; detection;
 KW quantitation; restorative therapy; polymorphic; ds.
 XX
 OS Homo sapiens.
 XX
 PN WO200140521-A2.
 XX
 PD 07-JUN-2001.
 XX
 PF 30-NOV-2000; 2000WO-US32758.
 XX
 PR 30-NOV-1999; 99US-0168138.
 PR 29-NOV-2000; 2000US-0726173.
 XX
 PA (CURA-) CURAGEN CORP.
 XX
 PI Shimkets RA, Leach M;
 XX
 DR WPI: 2001-356160/37.
 XX
 PT Polymorphic nucleic acid sequences, useful in genetic testing and
 PT therapy -
 XX
 PS Claim 1; Page 792; 2653pp; English.
 XX
 CC AAI73060 to AAI79867 represent isolated human polymorphic polynucleotide
 CC sequences (I), which contain single nucleotide polymorphisms (SNPs).
 CC AAI53114 to AAI53329 represent peptides related to human polymorphic
 CC polynucleotide sequences. The sequences can be used in gene and protein
 CC therapy, and in vaccine production. (I) and the polypeptides encoded by
 CC them may be used in the prevention, diagnosis and treatment of diseases
 CC associated with inappropriate expression of polymorphic polypeptides.
 CC For example, (I) may be used to treat disorders by rectifying mutations
 CC or deletions in a patient's genome that affect the activity of
 CC polypeptides by expressing inactive proteins or to supplement the
 CC patients own production of polypeptide. Additionally, (I) and its
 CC complementary sequences may also be used as DNA probes in diagnostic
 CC assays to detect and quantitate the presence of similar nucleic acids
 CC in samples, and therefore which patients may be in need of restorative
 CC therapy. The polypeptides encoded by (I) may be used as antigens in the
 CC production of antibodies specific for polymorphic polypeptides. The
 CC antibodies may also be used to down regulate expression and activity.
 CC The antibodies may also be used as diagnostic agents for detecting the
 CC presence of polymorphic polypeptides in samples.
 XX
 SQ Sequence 50 BP; 10 A; 17 C; 14 G; 9 T; 0 other;

alignment_scores:
 Quality: 43.00 Length: 9
 Ratio: 4.778 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 88.889
 alignment_block:
 US-09-836-410A-1 x AAI75479/rev ..
 Align seg 1/1 to reverse of: AAI75479 from: 1 to: 50

alignment_block:
US-09-836-410A-1 x AAL34450 ..
Align seg 1/1 to: AAL34450 from: 1 to: 50
6 LysIleTyrGluGluAlaTrpThrLysTyrProArgGlyLeuValProAr 22
|||||:|||||:|||||:|||||
3 AAATATATATGAGAAAGCTGGCAAC..... 26
22 glusLeuProLeuAsnPheLeuSerGlyGlu 32
|||||:|||||:|||||:|||||
27CCAGTGAACCTCTCGCGGAGAG 50

seq_name: /SIDS1/gcgdata/hold-geneseq/geneseq-emb1/NA1999.DAT: AAX88379

seq_documentation_block:
ID AAX88379 standard: DNA; 45 BP.

XX AAX88379;
AC AAX88379;
DT 30-SEP-1999 (first entry)
DE Antibody lambda light chain variable region PCR primer Rjambda0-B.
KW Haematopoietic; growth factor; PCR primer; mimetic; cell survival;
KW cell-proliferation; cell differentiation; cell activation; agonist;
KW growth factor inhibitor; nervous system cell; endodermal cell; therapy;
KW totipotent cell; embryonic stem cell; gene therapy; protection; allergy;
KW diagnostic; neutropenia; leukemia; aplastic anaemia; thrombocytopenia;
KW cancer cell; bone marrow transplant; myeloproliferative disease;
KW antibody; lambda light chain; variable region; ss.
XX Synthetic.
OS Synthetic.
XX WO9938008-A1.
XX PN
XX PD 29-JUL-1999.
XX PF 22-JAN-1999; 99WO-US01331.
XX PR 23-JAN-1998; 98US-0072253.
XX PS (PROL-) PROLIFARON INC.
XX PI Bowdish KS;
XX DR WPI; 1999-458732/38.

Identification of agonist or inhibitory antibodies to receptors that control cellular processes, used to modulate, e.g. proliferation of hematopoietic cells

Example 1; Page 115; 123pp; English.

This invention describes a novel method for identifying agonist or inhibitory antibodies (Ab) to receptors (R) involved in cell survival, proliferation, differentiation, or activations. The method is used to identify Ab that are growth factor mimetics and inhibitors and can regulate growth, differentiation, survival and activity of many different cell types, particularly hematopoietic cells (at various stages, of any lineage), but also nervous system cells, endodermal cells or totipotent (embryonic stem) cells. Ab are used (i) directly as therapeutic agents, e.g. for amplifying particular cell types, for ex vivo proliferation or differentiation of cells for use in gene therapy, to protect normal cells against chemotherapeutic agents, and to treat conditions such as allergy, (ii) as diagnostic/research reagents, e.g. for cell identification and sorting, (iii) to clone receptors and native factors that they mimic (also potential therapeutic agents). Typical therapeutic applications are in cases of neutropenia (of any etiology) or aplastic anemia, bone marrow transplants, myeloproliferative diseases (e.g. leukemia, thrombocytopenia or allergy) and inhibitory Ab. optionally coupled to a toxin, are used to kill cancer cells. The method is not subject to the usual limitations of monoclonal antibody technology

17 ArgGlyLeuValProArgLysLeuPro 25
|||||:|||||:|||||:|||||
43 AGAGGGCTGGTCCCGCTGCTGCTGCGG 17
seq_name: /SIDS1/gcgdata/hold-geneseq/geneseq-emb1/NA2001A.DAT: AAL34450
seq_documentation_block:
ID AAL34450 standard: DNA; 50 BP.
XX AAL34450;
XX DT 24-JAN-2002 (first entry)
XX DE Human SNP oligonucleotide #7658.
XX KW Immunosuppressive; immunostimulatory; antiinflammatory; cytostatic;
KW neuroprotective; antimicrobial; gene therapy; vaccine; amylase; cancer;
KW amyloid protein; angiopoietin; apoptosis related protein; cadherin;
KW cyclin; polymerase; oncogene; histone; kinase; colony stimulating factor;
KW complement related protein; cytochrome; kinesin; cytokine; interferon;
KW interleukin; G-protein coupled receptor; thioesterase; inflammation;
KW multifactorial disease; autoimmune disease; infection;
KW nervous system disease; ss.
XX Homo sapiens.
XX OS
XX PN WO200147944-A2.
XX PD 05-JUL-2001.
XX PF 28-DEC-2000; 2000WO-US35498.
XX PR 28-DEC-1999; 99US-0173419.
XX PS 27-DEC-2000; 2000US-0173419.
XX PI (CURA-) CURAGEN CORP.
XX PI Shimkets RA, Leach M;
XX DR WPI; 2001-465210/50.

Polymorphic nucleic acids encoding e.g. amylases, cyclins, polymerases, oncogenes and histones, useful for diagnosing and treating, e.g. cancer, autoimmune diseases and infections

Claim 1; Page 3600; 4143pp; English.

The present invention relates to oligonucleotides encoding polymorphic variants of proteins related to amylases, amyloid proteins, angiopoietin, apoptosis related proteins, cadherin, cyclin, polymerase, oncogenes, histones, kinases, colony stimulating factors, complement related proteins, cytochromes, kinesins, cytokines, interferons, interleukins, G-protein coupled receptors and thioesterases. The present sequence is one such oligonucleotide. The oligonucleotides and the peptides encoded by them may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate expression of the proteins listed above. Disorders that may be prevented, diagnosed and/or treated include multifactorial diseases with a genetic component, such as autoimmune diseases (e.g. rheumatoid arthritis, multiple sclerosis, diabetes, cancer, systemic lupus erythematosus and Grave's disease), inflammation, cancer (e.g. cancers of the bladder, brain, breast, colon and kidney, leukaemia), diseases of the nervous system and an infection of pathogenic organisms.

Sequence 50 BP; 18 A; 11 C; 13 G; 8 T; 0 other;

alignment_scores:
Quality: 42.50 Length: 27
Ratio: 2.833 Gaps: 1
Percent Similarity: 55.556 Percent Identity: 37.037

Claim 1; Page 551; 2745pp; English.

AAZ65654 to AAZ69578 represent human biallelic markers from the present invention, which contain a polymorphic base at position 24 of their nucleotide sequences. AAZ69579 to AAZ77440 represent amplification primers for the biallelic markers. The biallelic markers of the invention have a variety of uses: they can be used for high density mapping of the human genome, and in complex association studies and haplotyping studies which are useful in determining the genetic basis for disease states. Compositions and methods of the invention can also be useful for the identification of the targets for the development of pharmaceutical agents and diagnostic methods, as well as the characterisation of the differential efficacious responses to and side effects from pharmaceutical agents acting on a disease as well as other treatment.

N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297 and 3367, are not actually given a sequence in the Sequence Listing from the present invention.

Sequence 47 BP; 19 A; 5 C; 10 G; 13 T; 0 other;

alignment_scores:
Quality: 42.00 Length: 10
Ratio: 5.250 Gaps: 0
Percent Similarity: 80.000 Percent Identity: 70.000

alignment_block:

US-09-836-410A-1 x AAZ67157 ..

Align seg 1/1 to: AAZ67157 from: 1 to: 47

6 LysleTyrGluGluAlaTrpThrLysTyr 15
|||||
17 AAAATATATGACAGACAGTGGCCAAATAT 46

seq_name: /SIDS1/gcgdata/hold-geneseq/geneseq-embl/NA2001A.DAT:AAH75873

seq_documentation_block:
ID AAH75873 standard; DNA: 41 BP.

AAH75873;

26-OCT-2001 (first entry)

Human reverse transcriptase 13 coding sequence probe #1.

Human; reverse transcriptase 13; cytostatic; virucide; immunomodulatory; antiinflammatory; haemostatic; gene therapy; malignant tumour; haemopathy; HIV infection; immunological disease; inflammation; developmental disorder; probe; ss.

Homo sapiens.

WO200164893-A1.

07-SEP-2001.

26-FEB-2001; 2001WO-CN00280.

02-MAR-2000; 2000CN-0111806.

(BIOW-) BIOWINDOW GENE DEV INC SHANGHAI.

Mao Y, Xie Y;

WPI; 2001-550183/61.

New human reverse transcriptase 13 for diagnosing and treating developmental disorders, malignant tumor, hemopathy, human immunodeficiency virus infection, immunological diseases and inflammations

Example 7; Page 15; 34pp; Chinese.

The present invention relates to human reverse transcriptase 13 and its coding sequence (see AAH75868 and AAG66428). The reverse transcriptase and its coding sequence are useful in the diagnosis and treatment of malignant tumour, haemopathy, HIV infection, immunological diseases, various inflammations and developmental disorders. The present sequence is a probe, which was used in an example from the present invention.

Sequence 41 BP; 18 A; 7 C; 6 G; 10 T; 0 other;

alignment_scores:
Quality: 41.00 Length: 14
Ratio: 3.417 Gaps: 1
Percent Similarity: 85.714 Percent Identity: 42.857

alignment_block:

US-09-836-410A-1 x AAH75873/rev ..

Align seg 1/1 to reverse of: AAH75873 from: 1 to: 41

435 TrpLeuHisGluCysMetIleArgLeuPheHisSerValCys 448
|||||
39 TGGCTGCATAAATGCTCTT.....CTTTGAGAAATATCTGT 4

seq_name: /SIDS1/gcgdata/hold-geneseq/geneseq-embl/NA1999.DAT:AAH05908

seq_documentation_block:

ID AAX05908 standard; DNA: 47 BP.

AAX05908;

07-MAY-1999 (first entry)

Oligonucleotide probe meca945-29A18P.

Hybridization; RNase H; scissile linkage; nucleic acid detection; gene detection; polyamine; probe; DNA/RNA hybrid; ss.

Synthetic.

Key Location/Qualifiers

misc_RNA 14..17

/*tag= a

WO9901570-A2.

14-JAN-1999.

03-JUL-1998; 98WO-CA00631.

22-JUN-1998; 98US-0090273.

03-JUL-1997; 97US-0051827.

18-MAY-1998; 98US-0086021.

(IDBI-) ID BIOMEDICAL CORP.

Bryan RN, Cloney LP, Farnworth BA, Marostenmaki AJ;

WPI; 1999-106070/09.

Increasing the hybridization rate between two nucleic acids - using ribonuclease H (RNase H) and/or a polyamine, useful for detecting nucleic acids of interest in a sample

Example 1; Page 15; 45pp; English.

The invention relates to methods of increasing the hybridization rate between two nucleic acids. One method comprises construction of two nucleic acids and a polyamine, and hybridizing both nucleic acids together, under suitable conditions. Also provided is a similar method involving two nucleic acids with RNase H, where both nucleic acids do

CC effects from pharmaceutical agents acting on a disease as well as other
CC treatment.
CC N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297
CC and 3367, are not actually given a sequence in the Sequence Listing
CC from the present invention.

XX Sequence 47 BP; 20 A; 6 C; 10 G; 11 T; 0 other;

alignment_scores:
Quality: 41.00 Length: 15
Ratio: 4.556 Gaps: 0
Percent Similarity: 60.000 Percent Identity: 46.667

alignment_block:
US-09-836-410A-1 x AAZ67285/rev ..

Align seq 1/1 to reverse of: AAZ67285 from: 1 to: 47

97 ProProThrThrLeuLeuTrpValGlnTyrTyrLeuAlaGlnHis 111
||| ||| ::|||::|||::||| ||| ||| |||
47 CCGACAACCTTTATGCTGTGGATTTTACATTTTACATTTTACAC 3

seq_name: /SIDSI/gcgdata/hold-geneseq/geneseq-emb1/NA1989.DAT:AAAN92016

seq_documentation_block:
ID AAN92016 standard; DNA; 50 BP.

XX AC AAN92016;

XX DT 17-APR-1990 (first entry)

XX Sequence probe complementary to Neisseria gonorrhoeae genomic sequence
DE SSJK1 combined with the LIA2C amplifier sequence.

XX Neisseria gonorrhoeae genomic sequence SSJK1; LIA2C amplifier sequence;
KW file 'rcjk'; jkl.probes15(50).

XX OS Neisseria gonorrhoeae.

XX FH Key Location/Qualifiers

XX FT misc_feature 1..30

XX FT /*tag= a

XX FT /*sequence probe"

XX FT misc_feature 31..50

XX FT /*tag= b

XX FT /*LIA2C amplifier sequence"

XX PN W08903891-A.

XX PD 05-MAY-1989.

XX PF 14-OCT-1988; 88WO-US03644.

XX PR 30-SEP-1988; 88US-0252638, US-109282.

XX PA (CHIR-) CHIRON CORP.

XX PI Urdea MS, Warner B, Running JA, Kolberg JA, Clyne JM;

XX PI Sanchez-Pescador R;

XX DR WPI; 1989-150787/20.

XX Nucleic acid multimer for hybridisation assays

PT - having single-stranded oligo-nucleotide units

PT capable of binding specifically to sequences of interest.

XX Fig 14; : 112pp; English.

XX The sequence probe (tag a) is complementary to N. gonorrhoeae genomic

CC sequence SSJK1 from the file 'rcjk'. It is used to assay crude cellular

CC lysates and genomic DNA from different bacteria.. It is called

CC jkl.probes15(50).

XX SQ Sequence 50 BP; 11 A; 12 C; 13 G; 14 T; 0 other;

alignment_scores:
Quality: 41.00 Length: 13
Ratio: 4.100 Gaps: 0
Percent Similarity: 76.923 Percent Identity: 53.846

alignment_block:
US-09-836-410A-1 x AAN92016/rev ..

Align seq 1/1 to reverse of: AAN92016 from: 1 to: 50

519 GlySerLeuThrAsnArgAsnLeuGlnThrCysMetGlu 531
||| ::| |||::|||::||| ||| ||| |||
44 GGTCCTATGCTTAATCAGAATCTGCATATCTGCATGGAG 6

seq_name: /SIDSI/gcgdata/hold-geneseq/geneseq-emb1/NA2001A.DAT:AAAL34455

seq_documentation_block:

ID AAL34455 standard; DNA; 50 BP.

XX AC AAL34455;

XX DT 24-JAN-2002 (first entry)

XX DE Human SNP oligonucleotide #7663.

XX Immunosuppressive; immunostimulatory; antiinflammatory; cytostatic;
KW neuroprotective; antimicrobial; gene therapy; vaccine; amylase; cancer;
KW amyloid protein; angiotensin; apoptosis related protein; cadherin;
KW cyclin; polymerase; oncogene; histone; kinase; colony stimulating factor;
KW complement related protein; cytochrome; kinesin; cytokine; interferon;
KW interleukin; G-protein coupled receptor; thioesterase; inflammation;
KW multifactorial disease; autoimmune disease; infection;
KW nervous system disease; ss.

XX OS Homo sapiens.

XX PN W0200147944-A2.

XX PD 05-JUL-2001.

XX PF 28-DEC-2000; 2000WO-US35498.

XX PR 28-DEC-1999; 99US-0173419.

XX PR 27-DEC-2000; 2000US-0173419.

XX PA (CURA-) CURAGEN CORP.

XX PI Shimkets RA, Leach M;

XX DR WPI; 2001-465210/50.

XX Polymorphic nucleic acids encoding e.g. amylases, cyclins, polymerases,
PT oncogenes and histones, useful for diagnosing and treating, e.g.
PT cancer, autoimmune diseases and infections -
XX Claim 1; Page 3601; 4143pp; English.

The present invention relates to oligonucleotides encoding polymorphic
variants of proteins related to amylases, amyloid proteins, angiotensin,
apoptosis related proteins, cadherin, cyclin, polymerase, oncogenes,
histones, kinases, colony stimulating factors, complement related
proteins, cytochromes, kinesins, cytokines, interferons, interleukins,
G-protein coupled receptors and thioesterases. The present sequence is
one such oligonucleotide. The oligonucleotides and the peptides encoded
by them may be used in the prevention, diagnosis and treatment of
diseases associated with inappropriate expression of the proteins listed
above. Disorders that may be prevented, diagnosed and/or treated include
multifactorial diseases with a genetic component, such as autoimmune
diseases (e.g. rheumatoid arthritis, multiple sclerosis, diabetes,

CC systemic lupus erythromatosus and Grave's disease), inflammation, cancer
 CC (e.g. cancers of the bladder, brain, breast, colon and kidney,
 CC leukaemia), diseases of the nervous system and an infection of pathogenic
 CC organisms.
 XX

SQ Sequence 50 BP; 10 A; 13 C; 22 G; 5 T; 0 other;

alignment_scores:

Quality: 41.00 Length: 12
 Ratio: 4.100 Gaps: 0
 Percent Smilarity: 83.333 Percent Identity: 66.667

alignment_block:

US-09-836-410A-1 x AAL34455 ..

Align seg 1/1 to: AAL34455 from: 1 to: 50

364 GlyGlyProLysGluGluLeuIleProGluLysLeu 375

||||||| |||||||:|:|:|:|:|

14 GGAGGGCCGCCAGAGGAGTGCTGCGGAGACCTA 49

OM of: US-09-836-410A-1 to: EST:* out_format : pfs

Date: Jul 20, 2002 4:12 AM

About: Results were produced by the GenCore software, version 4.5,
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Command line parameters:

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-O=/sgn2_1/USPRO_pool/US09836410/runat_18072002_115032_29691/app_query.fasta_1.660  
-DB=EST -QFMT=fastcap -SUFFIX=p2nl5to50.rst -GAPOP=12.000  
-GAPEXT=4.000 -MINMATCH=0.100 -LOOPCL=0.000 -LOPEXT=0.000  
-QGAPOP=4.500 -QGAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500  
-FGAPOP=6.000 -FGAPEXT=7.000 -XGAPOP=10.000 -XGAPEXT=0.500  
-DELOP=6.000 -DELEXT=7.000 -START=1 -MATRIX=biosum62  
-TRANS=human40 cdi -LIST=45 -DOCALIGN=200 -THR_SCORE=pct  
-THR_MAX=100 -THR_MIN=0 -ALIGN=15 -MODE=LOCAL -OUTFMT=pfs  
-NORM_ext -HEAPSIZE=500 -MINLEN=15 -MAXLEN=50  
-USER=US09836410 -CGNL_1_2960 -NCPU=6 -ICPU=3 -LONGLOG  
-DEV_TIMEOUT=120 -WARN_TIMEOUT=30 -NO_XLPXY -WAIT -THREADS=1
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Search information block:

Query: US-09-836-410A-1

Query length: 593

Database: EST:*

Database sequences: 13736207

Database length: 1841457050

Search time (sec): 1532.560000

score_list:

Sequence	Strd	Orig	ZScore	EScore	Len	Documentation
gb_gss:A2774479	-	48.00	92.20	6.9e+04	33	A2774479 2M0004A05F Mouse 10kb F
gb_est1:AI657570	-	47.50	85.72	9.5e+04	40	AI657570 fc15c02.y1 Zebrafish W
gb_est1:AI232398	+	47.00	87.71	1.2e+05	46	AI232398 qx12h07.x1 NCI_CGAP_Ly
gb_est2:BF054800	+	45.00	83.94	2.0e+05	50	BF054800 7171g09.y1 NCI_CGAP_Br
gb_est1:AA590944	+	43.00	80.92	2.9e+05	50	AA590944 vm25f02.r1 Knowles Soli
gb_est1:AI813747	+	42.00	82.13	2.5e+05	37	AI813747 wk79a02.x1 NCI_CGAP_Pan
gb_est2:BM393347	-	41.00	78.09	4.2e+05	49	BM393347 50071-2-9-B10.f.1 Child
gb_est2:BM395447	-	41.00	78.09	4.2e+05	49	BM395447 50072-2-9-B10.f.1 Child
gb_est1:AI104189	+	41.00	77.91	4.3e+05	50	AI104189 AI104189 Sugano Homo sa
gb_est1:AI107491	+	41.00	77.91	4.3e+05	50	AI107491 AI107491 Sugano Homo sa
gb_gss:AZ846607	-	40.50	78.10	4.2e+05	45	AZ846607 1M0371N10F Mouse 10kb F
gb_gss:AZ860401	-	40.00	77.76	4.3e+05	43	AZ860401 2M0166E22F Mouse 10kb F
gb_est1:AI103783	+	40.00	77.55	4.5e+05	44	AI587842 AL587842 BP Chicken Bra
gb_est1:AI104282	+	40.00	76.40	5.2e+05	50	AI103783 AI103783 Sugano Homo sa
gb_est1:AI107535	+	40.00	76.40	5.2e+05	50	AI104282 AI104282 Sugano Homo sa
gb_est1:AA574989	-	39.50	76.20	5.3e+05	50	AA574989 vm34a03.r1 Knowles Soli
gb_gss:AZ582769	-	39.50	75.83	5.6e+05	47	AZ582769 1M0376B22F Mouse 10kb F
gb_est1:AI500599	+	39.50	75.83	5.6e+05	49	AI500599 tp93407.x1 NCI_CGAP_UC2
gb_est2:BJ043594	+	39.00	75.84	5.8e+05	45	BJ043594 BU043594 N18B Mochli nc
gb_est2:BI330882	+	39.00	75.45	5.8e+05	47	BI330882 602981270F1 NCI_CGAP_Ly
gb_est1:AI627881	+	39.00	75.07	6.1e+05	49	AI627881 AL627881 XGC-gastrula S
gb_est1:AA254893	+	39.00	74.89	6.3e+05	50	AA254893 mx78d07.r1 Soares mouse
gb_est1:AA966391	+	39.00	74.89	6.3e+05	50	AA966391 w4f01a1.r1 Aspergillus
gb_est1:AI107471	+	39.00	74.89	6.3e+05	50	AI107471 AI107471 Sugano Homo sa
gb_est1:AI107473	+	39.00	74.89	6.3e+05	50	AI107473 AI107473 Sugano Homo sa
gb_est1:AI107474	+	39.00	74.89	6.3e+05	50	AI107474 AI107474 Sugano Homo sa
gb_est1:AI107475	+	39.00	74.89	6.3e+05	50	AI107475 AI107475 Sugano Homo sa
gb_est1:AI107481	+	39.00	74.89	6.3e+05	50	AI107481 AI107481 Sugano Homo sa
gb_est1:AI107483	+	39.00	74.89	6.3e+05	50	AI107483 AI107483 Sugano Homo sa
gb_est1:AI107486	+	39.00	74.89	6.3e+05	50	AI107486 AI107486 Sugano Homo sa
gb_est1:AI107487	+	39.00	74.89	6.3e+05	50	AI107487 AI107487 Sugano Homo sa
gb_est1:AI107488	+	39.00	74.89	6.3e+05	50	AI107488 AI107488 Sugano Homo sa
gb_est1:AI107490	+	39.00	74.89	6.3e+05	50	AI107490 AI107490 Sugano Homo sa
gb_est1:AI107492	+	39.00	74.89	6.3e+05	50	AI107492 AI107492 Sugano Homo sa
gb_est1:AI107493	+	39.00	74.89	6.3e+05	50	AI107493 AI107493 Sugano Homo sa
gb_est1:AI107494	+	39.00	74.89	6.3e+05	50	AI107494 AI107494 Sugano Homo sa
gb_est1:AI107495	+	39.00	74.89	6.3e+05	50	AI107495 AI107495 Sugano Homo sa
gb_est1:AI107496	+	39.00	74.89	6.3e+05	50	AI107496 AI107496 Sugano Homo sa
gb_est1:AI107497	+	39.00	74.89	6.3e+05	50	AI107497 AI107497 Sugano Homo sa

gb_est1:AU107499 + 39.00 74.89 6.3e+05 50 ! AU107499 AU107499 Sugano Homo
gb_est1:AU107500 + 39.00 74.89 6.3e+05 50 ! AU107500 AU107500 Sugano Homo
gb_est1:AU107501 + 39.00 74.89 6.3e+05 50 ! AU107501 AU107501 Sugano Homo
gb_est1:AU107502 + 39.00 74.89 6.3e+05 50 ! AU107502 AU107502 Sugano Homo
gb_est1:AU107503 + 39.00 74.89 6.3e+05 50 ! AU107503 AU107503 Sugano Homo

seq_name: gb_gss:A2774479

seq_documentation_block:

LOCUS A2774479 33 bp DNA linear GSS 16-FEB-2001
DEFINITION 2M0004A05F Mouse 10kb plasmid UUGCLM library Mus musculus genomic
clone UUGC2M0004A05 F, DNA sequence.

ACCESSION A2774479.1 GI:12899972

VERSION GSS.

KEYWORDS house mouse.

SOURCE house mouse.

ORGANISM Mus musculus

Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 33)

AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,

Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly

M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.

and Wright,D., Weiss,R.

Mouse whole genome scaffolding with paired end reads from 10kb

plasmid inserts

Unpublished (2000)

Contact: Robert B. Weiss

University of Utah Genome Center

University of Utah

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT

84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

Plate: 0004 row: A column: 05

Seq primer: CGTGTGAACACGACGCCACG

Class: plasmid ends

High quality sequence step: 33.

FEATURES

Location/Qualifiers

1..33

/organism="Mus musculus"

/strain="C57BL/6J"

/db_xref="taxon:10090"

/clone="UUGC2M0004A05"

/sex="Male"

/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"

/note="Vector: PWD42nv; Purified genomic DNA from M.

musculus C57BL/6J (male) was obtained from the Jackson

Laboratory Mouse DNA Resource

(http://www.jax.org/resources/documents/dnares/). The DNA

was hydrodynamically sheared by repeated passage through a

0.005 inch orifice at constant velocity. The sheared DNA

was blunt end-repaired with T4 DNA polymerase and T4

polynucleotide kinase. Adaptor oligonucleotides were

ligated to the blunt ends in high molar excess. The

adaptor DNA was purified and size-selected for a 9.5 to

10.5 kb range using preparative agarose gel

electrophoresis. Vector DNA was prepared from a derivative

of pWD42 (g11473211419b/AP129072.1), a copy-number

inducible derivative of plasmid R1. The vector was ligated

with adaptors complementary to the insert adaptors and

purified. The sheared, adaptor mouse DNA was annealed to

adaptor vector DNA, and transformed into

chemically-competent E. coli XL10-Gold (Stratagene) cells

and selected for ampicillin resistance."

0 a 0 c 6 g 27 t

BASE COUNT

ORIGIN

alignment_scores:

analysis were selected following oligonucleotide hybridization fingerprinting of arrayed clones from zebrafish late somitogenesis (26 ss), adult liver or embryonic shield stage (5.6 h) libraries. Fingerprint data were used to computationally cluster cDNAs, and a single cDNA from each cluster was chosen for sequencing. In some cases multiple members of the same cluster were sequenced to assess clustering parameters or single clones were sequenced additional times to assess quality control."

BASE COUNT 8 a 11 c 14 g 7 t
ORIGIN

alignment_scores:
Quality: 47.50 Length: 11
Ratio: 4.750 Gaps: 1
Percent Similarity: 90.909 Percent Identity: 90.909

alignment_block:
US-09-836-410A-1 x AI657570/rev ..

Align seg 1/1 to reverse of: AI657570 from: 1 to: 40

560 ProTyAlaLeuAlaPheMetProGlyTyr 570
|||||
33 CCATATGCGGTACCC...ATGCCGCTGGGTAC 4

seq_name: gb_est1:AI223998

seq_documentation_block:
LOCUS AI223998 46 bp mRNA linear EST 21-DEC-1998
DEFINITION qx12h07.x1 NCI_CGAP_Lym12 Homo sapiens cDNA clone IMAGE:2001181 3' similar to TR:Q04154 Q04154 SALIVARY PROLINE-RICH PROTEIN RP15 PRECURSOR. ;contains element MER22 repetitive element ;, mRNA sequence.

ACCESSION AI223998
VERSION AI223998.1 GI:3806711
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 46)
REFERENCE NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
AUTHORS National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
JOURNAL: Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
unknown library type
Trace considered overall poor quality
Insert Length: 1534 Std Error: 0.00
Seq primer: -40UP from Gibco
High quality sequence stop: 1.
Location/Qualifiers
1. .46
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2001181"
/clone_lib="NCI_CGAP_Lym12"
/tissue_type="lymphoma, follicular mixed small and large cell"
/lab_host="DH10B"
/note="Organ: lymph node; Vector: pCMV-SPORT6; Site:1; SalI; Site:2; NotI; Cloned unidirectionally. Primer: Oligo dt. Average insert size 1.25 kb. Life Technologies catalog #: 11547-015"

BASE COUNT 19 a 17 c 10 g 0 t
ORIGIN

alignment_scores:

Quality: 48.00 Length: 10
Ratio: 4.800 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 80.000

alignment_block:
US-09-836-410A-1 x AZ774479/rev ..

Align seg 1/1 to reverse of: AZ774479 from: 1 to: 33

347 LysProGlnArgAsnProLysLysLysLys 356
|||||
32 AATCAACCAAAACCCCAAAAAA 3

seq_name: gb_est1:AI657570

seq_documentation_block:
LOCUS AI657570 40 bp mRNA linear EST 07-JUN-2001
DEFINITION fc15c02.y1 zebrafish Washu MPIMG EST Danio rerio cDNA clone IMAGE:3721442 5' similar to TR:O13017 O13017 WINGED HELIX TRANSCRIPTIONAL FACTOR MFH-1. ;, mRNA sequence.

ACCESSION AI657570
VERSION AI657570.1 GI:4755238
KEYWORDS EST.
SOURCE zebrafish.
ORGANISM Danio rerio
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes ; Cyprinidae; Danio.
1 (bases 1 to 40)
REFERENCE Clark,M., Johnson,S.L., Lehrach,H., Lee,R., Li,F., Marra,M., Eddy S., Hillier,L., Kucaba,T., Martin,J., Beck,C., Wylie,T., Underwood K., Steptoe,M., Theising,B., Allen,M., Bowers,Y., Person,B., Swaller,F., Gibbons,M., Pape,D., Harvey,N., Schurk,R., Ritter,E., Kohn,S., Shin,F., Jackson,Y., Cardenas,M., McCann,R., Waterston,R. and Wilson.R.
Washu Zebrafish EST Project 1998
Unpublished (1998)
Contact: Stephen L. Johnson
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1810
Email: zbrfish@watson.wustl.edu
cDNA Library Preparation: Matthew Clark. cDNA Library Arrayed by: Matthew Clark. DNA Sequencing by: Washington University Genome Sequencing Center Clone Distribution: Genome Systems, St. Louis, Missouri (web address: www.genomesystems.com) (email contact: info@genomesystems.com) and Research Genetics, Huntsville, Alabama (web address: www.resgen.com) (email contact: info@resgen.com) and RessourcenzentrumPrimatDatenbank, Berlin, Germany (web address: www.rzpd.de)
Trace considered overall poor quality
Possible reversed clone: similarity on wrong strand
Seq primer: T3 ET from Amersham
High quality sequence stop: 1.
Location/Qualifiers
1. .40
/organism="Danio rerio"
/db_xref="taxon:7955"
/clone="IMAGE:3721442"
/clone_lib="Zebrafish Washu MPIMG EST"
/sex="mixed"
/tissue_type="26 somite embryos, adult livers, shield stage embryos"
/lab_host="X1-blue MRF"
/note="Vector: pSPORT1; Site:1: NotI; Site:2: SalI; 1st strand cDNA was primed with a Not I oligo(dT)15 primer [5'pGACTACTTCTAGACGAGCGCGCCCTTTTITTTT3']; double-stranded cDNA was ligated to Sal I adaptors (BRL), digested with Not I and cloned into the Not I and Sal I sites of the pSPORT1 vector (BRL). Library was constructed by Matthew Clark (Lehrach lab; ICRF, London and Max Planck Institut fuer Molekulare Genetik, Berlin). cDNAs for EST

FEATURES
source
1. .40
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2001181"
/clone_lib="NCI_CGAP_Lym12"
/tissue_type="lymphoma, follicular mixed small and large cell"
/lab_host="DH10B"
/note="Organ: lymph node; Vector: pCMV-SPORT6; Site:1; SalI; Site:2; NotI; Cloned unidirectionally. Primer: Oligo dt. Average insert size 1.25 kb. Life Technologies catalog #: 11547-015"

BASE COUNT 19 a 17 c 10 g 0 t
ORIGIN

alignment_scores:

FEATURES
source
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/organism="Danio rerio"
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/clone_lib="Zebrafish Washu MPIMG EST"
/sex="mixed"
/tissue_type="26 somite embryos, adult livers, shield stage embryos"
/lab_host="X1-blue MRF"
/note="Vector: pSPORT1; Site:1: NotI; Site:2: SalI; 1st strand cDNA was primed with a Not I oligo(dT)15 primer [5'pGACTACTTCTAGACGAGCGCGCCCTTTTITTTT3']; double-stranded cDNA was ligated to Sal I adaptors (BRL), digested with Not I and cloned into the Not I and Sal I sites of the pSPORT1 vector (BRL). Library was constructed by Matthew Clark (Lehrach lab; ICRF, London and Max Planck Institut fuer Molekulare Genetik, Berlin). cDNAs for EST

FEATURES
source
1. .40
/organism="Danio rerio"
/db_xref="taxon:7955"
/clone="IMAGE:3721442"
/clone_lib="Zebrafish Washu MPIMG EST"
/sex="mixed"
/tissue_type="26 somite embryos, adult livers, shield stage embryos"
/lab_host="X1-blue MRF"
/note="Vector: pSPORT1; Site:1: NotI; Site:2: SalI; 1st strand cDNA was primed with a Not I oligo(dT)15 primer [5'pGACTACTTCTAGACGAGCGCGCCCTTTTITTTT3']; double-stranded cDNA was ligated to Sal I adaptors (BRL), digested with Not I and cloned into the Not I and Sal I sites of the pSPORT1 vector (BRL). Library was constructed by Matthew Clark (Lehrach lab; ICRF, London and Max Planck Institut fuer Molekulare Genetik, Berlin). cDNAs for EST

FEATURES
source
1. .40
/organism="Danio rerio"
/db_xref="taxon:7955"
/clone="IMAGE:3721442"
/clone_lib="Zebrafish Washu MPIMG EST"
/sex="mixed"
/tissue_type="26 somite embryos, adult livers, shield stage embryos"
/lab_host="X1-blue MRF"
/note="Vector: pSPORT1; Site:1: NotI; Site:2: SalI; 1st strand cDNA was primed with a Not I oligo(dT)15 primer [5'pGACTACTTCTAGACGAGCGCGCCCTTTTITTTT3']; double-stranded cDNA was ligated to Sal I adaptors (BRL), digested with Not I and cloned into the Not I and Sal I sites of the pSPORT1 vector (BRL). Library was constructed by Matthew Clark (Lehrach lab; ICRF, London and Max Planck Institut fuer Molekulare Genetik, Berlin). cDNAs for EST


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seq_name: gb_est1:A1813747

seq_documentation_block:
  A1813747
LOCUS       WK79a02.x1 NCI_CGAP_Panl Homo sapiens cDNA clone IMAGE:2421578 3'
DEFINITION similar to TR:Q39599 Q39599 EXTENSIN. ; contains element MSRI
repetitive element ;, mRNA sequence.
ACCESSION   A1813747
VERSION     A1813747.1 GI:5424962
KEYWORDS    EST.
SOURCE      human.
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1 (bases 1 to 37)
AUTHORS    NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE      National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor gene index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-re@mail.nih.gov
Life Technologies catalog #: 11548-013
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/dbrrp/image/image.html

Trace considered overall poor quality
Insert Length: 718 Std Error: 0.00
Seq primer: -40UP from Gibco
High quality sequence stop: 1.
Location/Qualifiers
1..37
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2421578"
/clone_lib="NCI_CGAP_Panl"
/tissue_type="adenocarcinoma"
/lab_host="DH10B"
/note="Organ: pancreas; Vector: pCMV-SPORT6; Site_1: SalI;
Site_2: NotI; Cloned unidirectionally. Primer: Oligo dt.
Average insert size 1.72 Kb. Life technologies catalog #:
11548-013"
BASE COUNT   19 a   16 c   1 g   1 t
ORIGIN

alignment_scores:
Quality: 42.00 Length: 9
Ratio: 4.667 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:
US-09-836-410A-1 x A1813747 ..

Align seg 1/1 to: A1813747 from: 1 to: 37

348 ProGlnArgAsnProlyslLysLysLys 356
|||||:::|||||||
6 CCCCAAAAAGCCCCAAAAA AAAA 32

seq_name: gb_est2:BM393347

seq_documentation_block:
LOCUS       BM393347
DEFINITION Tetrahymena thermophila cDNA (small fraction)
ACCESSION   BM393347
VERSION     BM393347.1 GI:18193400
KEYWORDS    EST.
SOURCE      Tetrahymena thermophila.
ORGANISM    Tetrahymena thermophila
Eukaryota; Alveolata; Ciliophora; Oligohymenophorea;
```


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annealed to

1

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seq_documentation_block:
LOCUS      AL587842      44 bp      mRNA      linear      EST 02-MAR-2001
DEFINITION      ROS064C05, BP Chicken Brain Library Gallus gallus cDNA clone
ACCESSION      AL587842
VERSION        AL587842.1 GI:13192876
KEYWORDS       EST.
SOURCE         chicken.
ORGANISM       Gallus gallus
               Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
               Archosauria; Aves; Neognathae; Galliformes; Phasianidae;
               1 (bases 1 to 44)
REFERENCE      Murray, F.
AUTHORS        BP Chicken Brain Library
JOURNAL        Unpublished (2001)
COMMENT        Contact: Frazer Murray
               Dept. Genomics and Bioinformatics
               Roslin Institute
               Roslin, Midlothian, EH25 9PS, UK
               Tel: +44 (0)131 527 4200
               Fax: +44 (0)131 440 0434
               Email: frazer.murray@bbsrc.ac.uk
               CGCGCCGCTTTTCTTTTCTTTTCTTTT 3' Poly A RNA purchased from Clonetech
               (*6854-
               Seq primer: M13F.
               Location/Qualifiers
                 1..44
                 /organism="Gallus gallus"
                 /db_xref="taxon:9031"
                 /clone="ROS064C05"
                 /clone_lib="BP Chicken Brain Library"
                 /tissue_type="Brain"
                 /dev_stage="Unknown"
                 /lab_host="DH10B"
                 /note="Vector: pSPOR1; Site_1: NotI; Site_2: SalI; Cloned
                 unidirectionally. Primer: Oligo dr. 5' adaptor sequence:
                 5' TCGACCTCGAG 3' ; 3' adaptor sequence: 5'
                 CGCGCCGCTTTTCTTTTCTTTTCTTTT 3' Poly A RNA purchased from
                 Clonetech (*6854-1)"
BASE COUNT      1 a      0 c      4 g      39 t
ORIGIN
alignment_scores:
  Quality: 40.00      Length: 12
  Ratio: 3.636      Gaps: 0
  Percent Similarity: 91.667      Percent Identity: 58.333
alignment_block:
  US-09-836-410A-1 x AL587842/rev ..
  Align seg 1/1 to reverse of: AL587842 from: 1 to: 44
  345 LysGLuLysProClnArgAsnProLysLysLysLys 356
  |||:||||| :|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
  42 AAAAAAAAAAAAAAAAAAAAAACCAAAAAAAAAAAAAA 7
seq_name: gb_est1:AU103783

seq_documentation_block:
LOCUS      AU103783      50 bp      mRNA      linear      EST 30-AUG-2001
DEFINITION      HEP14796, Sugano Homo sapiens cDNA library Homo sapiens cDNA clone
ACCESSION      AU103783
VERSION        AU103783.1 GI:13553304
KEYWORDS       EST.
SOURCE         human.
ORGANISM       Homo sapiens
               Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
               Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
               1 (bases 1 to 50)
REFERENCE
AUTHORS        Suzaki, Y., Taira, H., Tsunoda, T., Mizushima-Sugano, J., Sese, J., Hata
               , Y., Nakamura, Y., Suyama, A. and Sugano, S.
               Diverse transcriptional initiation revealed by fine, large-scale
               mapping of mRNA start sites
EMBO Rep. 2 (5), 388-393 (2001)
COMMENT        Contact: Yutaka Suzuki
               Department of Virology
               Institute of Medical Science, University of Tokyo
               4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
               Email: ysuzuki@ims.u-tokyo.ac.jp
               Suzaki, Y., Yoshitomo-Nakagawa, K., Maruyama, K., Suyama, A. and Sugano
               , S. Construction and characterization of a full length-enriched and
               a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).
FEATURES
  source
    1..50
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    /db_xref="taxon:9606"
    /clone="HEP14796"
    /clone_lib="Sugano Homo sapiens cDNA library"
BASE COUNT      10 a      13 c      11 g      16 t
ORIGIN
alignment_scores:
  Quality: 40.00      Length: 15
  Ratio: 4.000      Gaps: 0
  Percent Similarity: 66.667      Percent Identity: 53.333
alignment_block:
  US-09-836-410A-1 x AU103783/rev ..
  Align seg 1/1 to reverse of: AU103783 from: 1 to: 50
  103 TrpValGlnTyrTyrLeuAlaGlnHisTyrAspLysIleGlyCln 117
  |||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
  49 TGGCTACTCTCCTTGGCGGAGGAGCGACAACTTGGACAA 5
seq_name: gb_est1:AU104282

seq_documentation_block:
LOCUS      AU104282      50 bp      mRNA      linear      EST 30-AUG-2001
DEFINITION      HEP05565, Sugano Homo sapiens cDNA library Homo sapiens cDNA clone
ACCESSION      AU104282
VERSION        AU104282.1 GI:13553803
KEYWORDS       EST.
SOURCE         human.
ORGANISM       Homo sapiens
               Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
               Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
               1 (bases 1 to 50)
REFERENCE
AUTHORS        Suzaki, Y., Taira, H., Tsunoda, T., Mizushima-Sugano, J., Sese, J., Hata
               , Y., Nakamura, Y., Suyama, A. and Sugano, S.
               Diverse transcriptional initiation revealed by fine, large-scale
               mapping of mRNA start sites
EMBO Rep. 2 (5), 388-393 (2001)
COMMENT        Contact: Yutaka Suzuki
               Department of Virology
               Institute of Medical Science, University of Tokyo
               4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
               Email: ysuzuki@ims.u-tokyo.ac.jp
               Suzaki, Y., Yoshitomo-Nakagawa, K., Maruyama, K., Suyama, A. and Sugano
               , S. Construction and characterization of a full length-enriched and
               a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).
FEATURES
  source
    1..50
    /organism="Homo sapiens"
    /db_xref="taxon:9606"
    /clone="HEP05565"

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AUTHORS        Suzaki, Y., Taira, H., Tsunoda, T., Mizushima-Sugano, J., Sese, J., Hata
               , H., Ota, T., Isogai, T., Tanaka, T., Morishita, S., Okubo, K., Sakaki
               , Y., Nakamura, Y., Suyama, A. and Sugano, S.
               Diverse transcriptional initiation revealed by fine, large-scale
               mapping of mRNA start sites
EMBO Rep. 2 (5), 388-393 (2001)
COMMENT        Contact: Yutaka Suzuki
               Department of Virology
               Institute of Medical Science, University of Tokyo
               4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
               Email: ysuzuki@ims.u-tokyo.ac.jp
               Suzaki, Y., Yoshitomo-Nakagawa, K., Maruyama, K., Suyama, A. and Sugano
               , S. Construction and characterization of a full length-enriched and
               a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).
FEATURES
  source
    1..50
    /organism="Homo sapiens"
    /db_xref="taxon:9606"
    /clone="HEP14796"
    /clone_lib="Sugano Homo sapiens cDNA library"
BASE COUNT      10 a      13 c      11 g      16 t
ORIGIN
alignment_scores:
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DEFINITION      HEP05565, Sugano Homo sapiens cDNA library Homo sapiens cDNA clone
ACCESSION      AU104282
VERSION        AU104282.1 GI:13553803
KEYWORDS       EST.
SOURCE         human.
ORGANISM       Homo sapiens
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               Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
               1 (bases 1 to 50)
REFERENCE
AUTHORS        Suzaki, Y., Taira, H., Tsunoda, T., Mizushima-Sugano, J., Sese, J., Hata
               , Y., Nakamura, Y., Suyama, A. and Sugano, S.
               Diverse transcriptional initiation revealed by fine, large-scale
               mapping of mRNA start sites
EMBO Rep. 2 (5), 388-393 (2001)
COMMENT        Contact: Yutaka Suzuki
               Department of Virology
               Institute of Medical Science, University of Tokyo
               4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
               Email: ysuzuki@ims.u-tokyo.ac.jp
               Suzaki, Y., Yoshitomo-Nakagawa, K., Maruyama, K., Suyama, A. and Sugano
               , S. Construction and characterization of a full length-enriched and
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us-09-836-410a-1.p2n15to50.rst

Mon Jul 22 09:40:56 2002

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XX      WO200179506-A2.
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XX      25-OCT-2001.
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XX      17-APR-2001; 2001WO-US12548.
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XX      17-APR-2000; 2000US-197977P.
XX      (CHIL-) CHILDRENS HOSPITAL RES FOUND.
XX
XX      Gendron RL, Paradis H;
XX
XX      WPI; 2002-026032/03.
XX      P-PSDB; AAE13589, AAE13590, AAE13591, AAE13592.
XX
XX      Novel tubedown-1 protein comprising anti-angiogenic activity is useful
XX      for treating angiogenesis-associated disease related to ocular
XX      neovascularization, e.g., diabetic retinopathy, retinopathy of
XX      prematurity.
XX
XX      Claim 10; Page 56-58; 85pp; English.
XX
XX      The present invention relates to tubedown-1 (tbdn-1) proteins and
XX      their corresponding cDNAs. Tbdn-1 proteins having anti-angiogenic
XX      activity are associated with acetyl transferase activity. They
XX      regulate endothelial differentiation through protein acetylation,
XX      DNA-binding or by interacting with and/or acetylating other protein,
XX

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Mon Jul 22 09:40:54 2002

CC targets important for endothelial differentiation. In normal adult
 CC eyes, tbdn-1 is expressed highly in the corneal endothelium proper
 CC and in the vascular endothelium of the limbus and retina. Tbdn-1
 CC proteins are useful for preventing, treating, inhibiting or delaying
 CC the onset of angiogenesis-associated disease, related to ocular
 CC neovascularisation, e.g., a retinal disease, such as preferably
 CC diabetic retinopathy or retinopathy of prematurity, or primary
 CC hyperplastic vitreous, macular degeneration and any other conditions
 CC involving ocular neovascularisation. Tbdn-1 proteins are also useful
 CC for treating any pathological neovascularisation condition such as
 CC head trauma, spinal trauma, stroke, haemorrhagic shock, arthritis,
 CC arteriosclerosis, angiofibroma, delayed wound healing, granulations,
 CC cancer, burns, scars, nonunion fractures, retrolental fibroplasia,
 CC solid tumour growth. Proteins of the invention are also useful for
 CC treating ocular neovascularisation conditions such as chronic glaucoma,
 CC sickle cell retinopathy, corneal neovascularisation, rubeosis iritis,
 CC uveitis, neovascularisation of the optic nerve. Sequences of the
 CC invention are also used in gene therapy. The present sequence is
 CC a cDNA encoding tubedown-1 proteins.

XX Sequence 3418 BP; 1157 A; 604 C; 704 G; 953 T; 0 other;

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AC AAH77156;

DT 21-JAN-2002 (first entry)

DE Human tubedown-1 cDNA.

KW Human; tubedown-1; tbdn-1; antisense; cytostatic; osteopathic;
 KW bone tumour; osteosarcoma; Ewings sarcoma; metastasis; ss.

OS Homo sapiens.

FH Key Location/Qualifiers
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XX WO200179505-A2.

PN 25-OCT-2001.

PD 17-APR-2001; 2001WO-US12435.

PE 17-APR-2000; 2000US-197977P.

PR 17-APR-2001; 2001US-0836410.

XX (CHIL-) CHILDRENS HOSPITAL RES FOUND.

PI Gendron RL, Paradis H;

XX WPI; 2002-017618/02.

DR P-PSDB; AAG77907.

XX Nucleic acid molecules antisense to the tubedown-1 gene prevent
 PT overexpression of tubedown-1 protein and are useful to treat
 PT osteosarcoma and Ewing's Sarcoma family of tumours

XX Claim 1; Page 36-38; 56pp; English.

XX

CC The sequence represents a new human gene, tubedown-1 (tbdn-1). The
 CC invention relates to a novel isolated nucleic acid of the tubedown-1
 CC gene, and antisense nucleic acids to tbdn-1. The polynucleotides and
 CC protein of the invention have cytostatic and osteopathic activity. The
 CC polynucleotides of the invention may be used in antisense-therapy/gene
 CC therapy. They are useful in the treatment of bone tumours, especially
 CC osteosarcoma and Ewings sarcoma family of tumours. The compounds of the
 CC invention may also be useful for the prevention of metastases from these
 CC types of tumours, either alone or in combination with radiotherapy and/or
 CC chemotherapeutic agents.
 XX
 SQ Sequence 3418 BP; 1157 A; 604 C; 704 G; 953 T; 0 other;

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 Ratio: 5.234 Gaps: 0

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XX AC AAH77158;

XX DT 21-JAN-2002 (first entry)

XX DE Human tubedown-1 base pairs 3418-1 antisense cDNA.

XX KW Human; tubedown-1; tbdn-1; antisense; cytostatic; osteopathic;

XX KW bone tumour; osteosarcoma; Ewings sarcoma; metastasis; ss.

XX OS Homo sapiens.

XX PN WO200179505-A2.

XX PD 25-OCT-2001.

XX PF 17-APR-2001; 2001WO-US12435.

XX PR 17-APR-2000; 2000US-197977P.

XX PR 17-APR-2001; 2001US-0836410.

XX PA (CHIL-) CHILDRENS HOSPITAL RES FOUND.

XX PI Gendron RL, Paradis H;

XX DR WPI; 2002-017618/02.

XX PT Nucleic acid molecules antisense to the tubedown-1 gene prevent

XX PT overexpression of tubedown-1 protein and are useful to treat

XX PT osteosarcoma and Ewing's Sarcoma family of tumours

XX PS Claim 7; Page 39-41; 56pp; English.

XX PS The sequence represents tubedown-1 (tbdn-1) bases 3418-1 antisense cDNA.

XX CC The invention relates to a novel isolated nucleic acid of the tubedown-1

XX CC gene, and antisense nucleic acids to tbdn-1. The polynucleotides and

XX CC protein of the invention have cytostatic and osteopathic activity. The

XX CC polynucleotides of the invention may be used in antisense-therapy/gene

XX CC therapy. They are useful in the treatment of bone tumours, especially

XX CC osteosarcoma and Ewings sarcoma family of tumours. The compounds of the

XX CC invention may also be useful for the prevention of metastases from these

XX CC types of tumours, either alone or in combination with radiotherapy and/or

XX CC chemotherapeutic agents.

XX SQ Sequence 3418 BP; 953 A; 704 C; 604 G; 1157 T; 0 other;

alignment_scores:
Quality: 3104.00 Length: 593
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2161 AAGCAGCGAGAATTGCTATTGAGATCTATTTGAAGCTTCATGACAACCT 2112
301 LeuThrAspGluAsnLysGluHisGluAlaAspThrAlaAsnMetSerAs 317
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2111 CTGACAGATGAGAACAAAGAACCCAGGCTGATACAGCAACATGTCTGA 2062
317 pLysGluLeuLysLysLeuArgAsnLysGlnArgAlaGlnLysLysA 334
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2061 CAAAGAGCTAAAGAAATGCGTAAATAACAAAGAAAGAGCTCAAAAGAAAG 2012
334 lAsGlnIleGluGluLysLysAsnAlaGluLysGluLysProGlnArg 350
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2011 CCCAGATTGAAGAGAGAAAAAATGCCGAAAAAGAAAGCCGCAACGG 1962
351 AsnProLysLysLysAspAspAspGluGluIleGlyGlyProLy 367
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1961 AATCCGAAAAAGAAAGAGATGATGATGACGAAAGAAATGGAGGCCCAA 1912
367 sGluGluLeuLeuProGluLysLeuAlaLysValGluThrProLeuGluG 384
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1911 AGAAGAGCTTATCCCTGAGAACTGCCCAAGGTTGAAACTCCATTGGAAG 1862
384 luAlaIleLysPheLeuThrProLeuLysAsnLeuValLysAsnLysIle 400
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1861 AAGCTATTAAAGTTTTTAACACCATTTGAAGAACTTGGTGAAGAAAGATA 1812
401 GluThrHisLeuPheAlaPheGluIleTyrPheArgLysGluLysPheLe 417
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1811 GAAACTCATCTTTTGCCTTTGAGATCTACTTTAGGAAAGAAAGTTTCT 1762
417 uLeuMetLeuGlnSerValLysArgAlaPheAlaIleAspSerSerHisP 434
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1761 TTTGATGCTACAATCAGTAAAGCGGCGCTTTGCTATTGATTCTAGTCATC 1712
434 rOTpLeuHisGluCysMetIleArgLeuPheHisSerValCysGluSer 450
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1711 CCTGGCTTCATGAGTGCATGATTCGACTCTTTTCATCTCTGTGTGGAAGT 1662
451 LysAspLeuProGluThrValArgThrValLeuLysGlnGluMetAsnAr 467
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1661 AAGACTTACCAGAAACAGATTAGAACAGTATTAAACAAAGAAATGAATCG 1612
467 gLeuPheGlyAlaThrAsnProLysAsnPheAsnGluThrPheLeuLysA 484
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1611 TCCTTTTGGAGCAACAATCCAAAGAAATTTTAATGAAACCTTTCTGAAAA 1562
484 rGAsnSerAspSerLeuProHisArgLeuSerAlaAlaLysMetValTyr 500
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1561 GGAATTCGATTCTATTCGCCCATAGATTATCAGCTGCCAAATGGTATAT 1512
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1511 TATTAGATTCTTCTAGTCAAAAACAGCAATAGAGCTGGCGCAACACT 1462
517 uAspGlySerLeuThrAsnArgAsnLeuGlnThrCysMetGluValLeuG 534
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1461 TGATGGATCCCTCACCACCAAGAAACCTTCAGACTTGCATGGAAGTGTGG 1412
534 luAlaLeuCysAspGlySerLeuArgAspCysLysGluAlaAlaGluAla 550
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1411 AAGCCTTGTGTGATGGTAGCCTACGAGACTGTAAAGAAAGCTGCCGAAGCC 1362
551 TyrArgAlaSerCysHisLysLeuPheProTyrAlaLeuAlaPheMetPr 567
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 17 gGlyLeuValProArgLysLeuProLeuAsnPheLeuSerGlyGluLysP 34
 585 GGGACTGGTCCCAAGAAGCTGCCGTTAAACTTTTATCTGGTGAAGT 634
 34 helysGluCysLeuAspArgPheLeuArgMetAsnPheSerLysGlyCys 50
 635 TTAAGAATGTTGGATAAGTTCTTAAGGATGAATTCAGCAAGGGTTC 684
 51 ProProValPheAsnThrLeuArgSerLeu.TyrArgAspLysGluLys. 66
 685 CCACCACTCTTCAATACTTTAAGATCATTTACTACCAAGACAAAGAAAGT 734
 67 ValAlaIleValGluLeuValValGlyTyrGluThrSerLeuLysSe 83
 735 GTGCAATCATGAAAGTTAGTAGTAGTTATGAAACCTCTCTAAAGAG 784
 83 rCysArgLeuPheAsnProAsnAspAspGlyLysGluGluProProThrT 100
 785 CTGCCGCTATTTAACCCCAATGATGAGAAAGGAGGAGAACCAACCA 834
 100 hrLeuLeuTyrValGlnTyrTyrLeuAlaGlnHisTyrAspLysIleGly 116
 835 CATTACTTTGGGTCCAGTACTTGTGCACACATTTATGACAAATTTGT 884
 117 GlnProSerIleAlaLeuGluTyrIleAsnThrAlaIleGluSerThrPr 133
 885 CAGCCATCTATTGCTTTGGAGTACATAAATACTGCTATTGAAAGTACAC 934
 133 oThrLeuIleGluLeuPheLeuValLysAlaLysIleTyrLysHisAla 150
 935 TACATTATAGAACTCTTTCTGCGAAAGCTAAATCTATAAGCATGCTG 984
 150 lyAsnIleLysGluAlaAlaArgTrpMetAspGluAlaGlnAlaLeuAsp 166
 985 GAAATATTAAGAAGCTGCAAGGTGGATGGATGAGCCCGCTTGGAC 1034
 167 ThrAlaAspArgPheIleAsnSerLysCysAlaLysTyrMetLeuLysAl 183
 1035 ACAGCAGACAGATTATCAACTCCCAATGTGCAAAATACATGCTAAAGC 1084
 183 aAsnLeuIleLysGluAlaGluGluMetCysSerLysPheThrArgGlu 200
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 250 lnPheAspPheHisThrTyrCysMetArgLysIleThrLeuArgSerTyr 266
 1285 AGTTTGACTTTTCATACATCTGATGAGGAAGATTACCTTAGATCATAT 1334

567 oProGlyTyrGluGluAspMetLysIleThrValAsnGlyAspSerSera 584
 1311 TCCTGGATACGAAGAGGATATGAAGATCACAGTGAACGGAGATAGTTCTG 1262

584 laGluThrGluGluLeuAlaAsnGluIle 593
 1261 CAGAAACGGAAGAACTGGCCATGAATC 1233

seq_name: /SIDS1/gcgdata/hold-geneseq/geneseqn-emb1/NA2001B.DAT:AAS71925

seq_documentation_block:

ID AAS71925 standard; cDNA; 2477 BP.

XX AC AAS71925;

XX DT 13-FEB-2002 (first entry)

XX DE DNA encoding novel human diagnostic protein #7729.

XX KW Human; chromosome mapping; gene mapping; gene therapy; forensic;

XX KW food supplement; medical imaging; diagnostic; genetic disorder; ss.

XX OS Homo sapiens.

XX PN WO200175067-A2.

XX PD 11-OCT-2001.

XX PF 30-MAR-2001; 2001WO-US08631.

XX PR 31-MAR-2000; 2000US-0540217.

XX PR 23-AUG-2000; 2000US-0649167.

XX PA (HYSE-) HYSEQ INC.

XX PI Drmanac RT, Liu C, Tang YT;

XX WPI; 2001-639362/73.

XX DR P-PSDB; ABG07738.

XX PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -

XX PS Claim 1; SEQ ID NO 7729; 103pp; English.

XX CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.

XX CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.

XX CC Sequence 2477 BP; 859 A; 437 C; 522 G; 659 T; 0 other;

XX SQ

267 ValAspLeuLeuLysLeuGluAspValLeuArgGlnHisProPheTyrPh 283
1335 GTGGACTTATTAAACTAGAGATGTACTTCGACAGCATCCTATTTCCT 1384
283 eLysAla...AlaArgIleAlaIleGluIleTyrLeuLysLeuHisAspA 299
1385 CAAGGCGAGGCAAGAAATTCCTATAGATCTATTGGAAGGCTTCATGACA 1434
299 snProLeuThrAspGluAsn...LysGluHis.GluAla.AsphThrAla. 313
1435 ACCCCCTTACAGATGAGGAATTAAGGAACACGAGGCTGGATACAGCCA 1484
314 AsnMetSerAspLysGluLeuLysLysLeuArgAsnLysGlnArgAl 330
1485 AACATGCTGACAAAGAGCTAAAGAACCTACGTAATAAACAAGAACAGC 1534
330 aGlnLysLysAlaGlnIleGluGluLysLysLysAsnAlaGluLysGlu 347
1535 TCAAAAGAAAGCCAGATGAAGAAGAGAAAGAAATGCCAGAAAAGAAA 1584
347 ysProGlnArgAsnProLysLysLysLysAspAspAspGluGluIle 363
1585 AGCAGCAGAGAAATCAGAAAAGAGAGAGATGATGATGAGGAGATA 1634
364 GlyClyProLysGluGluLeuIleProGluLysLeuAlaLysValGluTh 380
1635 GGAGGTCCAAAGAAAGAACTATTCCAGAGAAAGTGGCCAAAGTTGAAC 1684
380 rProLeuGluAlaIleLysPheLeuThrProLeuLysAsnLeuVal 397
1685 TCCATTGGAAGAGCTATTAAATTTTAAACCCGTTGAAGAACTTGTGA 1734
397 ysAsnLysIleGluThrHisLeuPheAlaPheGluIleTyrPheArgLys 413
1735 AGAACAAGATAGAGACTCATCTTTTCCCTTCGAGATTACTTTAGGAAA 1784
414 GluLysPheLeuLeuMetLeuGlnSerValLysArgAlaPheAlaIleAs 430
1784 1784
430 pSerSerHisProTrpLeuHisGluCysMetIleArgLeuPheHisSerV 447
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447 alCysGluSerLysAspLeuProGluThrValArgThrValLeuLysGln 463
1784 1784
464 GluMetAsnArgLeuPheGlyAlaThrAsnProLysAsnPheAsnGluTh 480
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1785GCTGCCA 1791
497 ysMetValTyrTyrLeuAspSerSerGlnLysArgAlaIleGluLeu 513
1792 AAATGGTATTACTATAGATCTCTTCTAGTCAGAAAGCGAGCTATAGATTG 1841
514 AlaThrThrLeuAspGlySerLeuThrAsnArgAsnLeuGlnThrCysMe 530
1842 GCAACAACACTTGATGAATCTCTACTACAGAAACCTCCAGACATGTAT 1891
530 tGluValLeuGluAlaLeuCysAspGlySerLeuArgAspCysLysGluA 547
1892 GGAGGTATTGGAAGCCTTGTATGATGGTAGCCTAGGAGACTGTAAGAGAG 1941
547 IaAlaGluAlaTyrArgAlaSerCysHisLysLeuPheProTyrAla. 562
1942 CTGCTGAAATTTATAGACAAATTCCTATAGCTTTACCCCTTTATGCT 1991

563 LeuAlaPhe...MetProProGlyTyrGluGluAspMetLysIle.ThrV 578
1992 TTGGCCTTTTCATGCCCGCCCGGATATGAAGAGATATGAAGATCCACAG 2041
578 alAsnGlyAspSerSerAlaGluThrGluGluLeuAla.AsnGluIle 593
2042 TTAATGGAGATAGTTCTCGAAGCTGGAAGAACTGGCCCAATGAAATT 2090

seq_name: /SIDS1/gcgdata/hold-geneseq/geneseqn-emb1/NA2002.DAT:AAH77157

seq_documentation_block:

ID AAH77157 standard; cDNA; 1413 BP.

XX AAH77157;

AC AAH77157;

XX 21-JAN-2002 (first entry)

XX Human tubedown-1 base pairs 1413-1 antisense cDNA.

XX Human; tubedown-1; tbdn-1; antisense; cytostatic; osteopathic;
bone tumour; osteosarcoma; Ewings sarcoma; metastasis; ss.

XX Homo sapiens.

XX WO200179505-A2.

XX 25-OCT-2001.

XX 17-APR-2001; 2001WO-US12435.

XX 17-APR-2000; 2000US-197977P.

XX 17-APR-2001; 2001US-0836410.

XX (CHIL-) CHILDRENS HOSPITAL RES FOUND.

XX Gendron RL, Paradis H;

XX WPI; 2002-017618/02.

XX Nucleic acid molecules antisense to the tubedown-1 gene prevent
overexpression of tubedown-1 protein and are useful to treat
osteosarcoma and Ewing's Sarcoma family of tumours

XX Claim 6; Page 38-39; 56pp; English.

XX The sequence represents tubedown-1 (tbdn-1) bases 1413-1 antisense cDNA.
The invention relates to a novel isolated nucleic acid of the tubedown-1
gene; and antisense nucleic acids to tbdn-1. The polynucleotides and
protein of the invention have cytostatic and osteopathic activity. The
polynucleotides of the invention may be used in antisense-therapy/gene
therapy. They are useful in the treatment of bone tumours, especially
osteosarcoma and Ewings sarcoma family of tumours. The compounds of
invention may also be useful for the prevention of metastases from these
types of tumours, either alone or in combination with radiotherapy and/or
chemotherapeutic agents.

XX Sequence 1413 BP; 359 A; 314 C; 244 G; 496 T; 0 other;

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Percent Identity: 100.000

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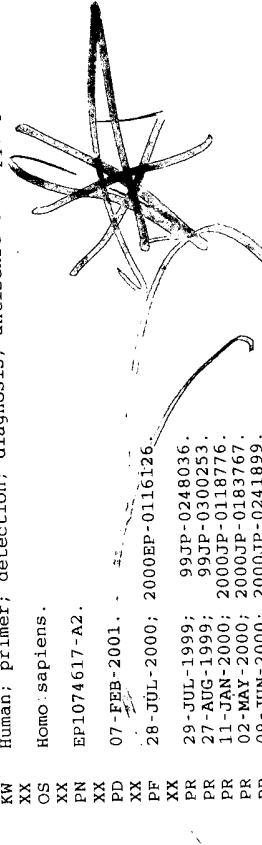
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17 gGlyLeuValProArgLysLeuProLeuAsnPheLeuSerGlyGluLysP 34
1956 GGGACTCGTCCCAAGAAAGCTCCCTTAACTTTATCTGGAGAGAGT 907
34 helysGluCysLeuAspArgPheLeuArgMetAsnPheSerLysGlyCys 50
906 TTAAGGAGTGTTGGATAGTTCCTTAAAGATGAATTCAGCAAGGCTGT 857
51 ProProValPheAsnThrLeuArgSerLeuTyrArgAspLysGluLysVa 67
856 CCACCTGTCTTCAATACCTTGGTCTTTATACAGAGATAAGAGAGGT 807
67 lAlalleValGluLeuValValGlyTyrGluThrSerLeuLysSerC 84
806 GGCAATCGTAGAAGAACTAGTGTGTTTAAAGAACTTCTTAAAAAGTT 757
84 ysArgLeuPheAsnProAsnAspAspGlyLysGluGluProProThrThr 100
756 GTGCCCTATTAAACCCCAATGATGATGGAAGGAGGAACCTCCAACACA 707
101 LeuLeuTrpValGlnTyrTyrLeuAlaGlnHisTyrAspLysIleGlyG 117
706 TTACTTTGGGTCCAGTACTATTTGGCACACATTTATGATAAAATTTGTCA 657
117 nProSerIleAlaLeuGluTyrIleAsnThrAlaIleGluSerThrProt 134
656 GCCATCCATTGCTCTGGAAATACATAAATACTGCAATTTGAAAGTACACCA 607
134 hrLeuIleGluLeuPheLeuValLysAlaLysIleTyrLysHisAlaGly 150
606 CATTGATAGAACTCTCTTCTTAAAGAGCTAAATCTATAAGCATGCTGGG 557
151 AsnIleLysGluAlaAlaArgTrpMetAspGluAlaGlnAlaLeuAspTh 167
556 AATATTAAAGAAAGCTGCAGGTGATGATGATGATGATGATGATGATGAT 507
167 rAlaAspArgPheIleAsnSerLysCysAlaLysTyrMetLeuLysAla 184
506 AGCAGACAGATTATTATTTCCAGTGTGCAAAATACATGTTAAAGGCA 457
184 snLeuIleLysGluAlaGluMetCysSerLysPheThrArgGluGly 200
456 ACCTGATTAAAGAGAGCTGAAGAAATGTTTCCAGTTTACGAGGAAGA 407
201 ThrSerAlaValGluAsnLeuAlaGlnMetGlnCysMetTrpPheGlnTh 217
406 ACTTCAGCGGTAGAGAACTGATGAATGCAATGCTATGTGTTCACAC 357
217 rGluCysAlaGlnAlaTyrLysAlaMetAsnLysPheGlyGluAlaLeu 234
356 AGAGTGTGCTCAGGCATACAAAGCAATGAACAAATTTGTGTAAGCACTTA 307
234 ysLysCysHisGluIleGluArgHisPheIleGluIleThrAspAspGln 250
306 AGAAATGTCATGAATTTGAGAGACATTTATAGAAATCACCAGTACCCAG 257
251 PheAspPheHisThrTyrCysMetArgLysIleThrLeuArgSerTyrVa 267
256 TTTGACTTTTATACATACATGATGATGAGGAAGATCACCCCTTAGATCATGT 207
267 lAspLeuLeuLysLeuGluAspValLeuArgGlnHisProPheTyrPheL 284
206 GGACTTTATTAATACTAGAGATGCTCTGACAGCATCCATTTTACTTCA 157
284 ysAlaAlaArgIleAlaIleGluIleTyrLeuLysLeuHisAspAsnPro 300
156 AAGCAGCGGAAGTGTCTTTGAGATCTATTGGAAGCTTCATGACAAACCC 107
301 LeuThrAspGluAsnLysGluHisGluAlaAspThrAlaAsnMetSerAs 317
106 CTGACAGATGAGAACAAAGAACACGAGGCTGATACAGCAACATGCTGA 57
317 pLysGluLeuLysLysLeuArgAsnLysGlnArgAlaGlnLysLysA 334

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seq_documentation_block:
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AC AAH16408;
XX 26-JUN-2001 (first entry)
DT
XX Human cDNA sequence SEQ ID NO:15380.
DE
XX Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.
XX Homo.sapiens.
XX EP1074617-A2.
XX
PD 07-FEB-2001.
XX 28-JUL-2000; 2000EP-0116126.
PF
XX 29-JUL-1999; 99JP-0248036.
PR 27-AUG-1999; 99JP-0300253.
PR 11-JAN-2000; 2000JP-0118776.
PR 02-MAY-2000; 2000JP-0183767.
PR 09-JUN-2000; 2000JP-0241899.
XX
PA (HELI-) HELIX RES INST.
XX Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX WPI; 2001-318749/34.



Primer sets for synthesizing polynucleotides, particularly the 5602 full-length cDNAs defined in the specification, and for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs -
Claim 8; SEQ ID 15380; 2537bp + CD ROM; English.
The present invention describes primer sets for synthesizing 5602 full-length cDNAs defined in the specification. Where a primer set comprises: (a) an oligo-dT primer and an oligonucleotide complementary to the complementary strand of a polynucleotide which comprises one of the 5602 nucleotide sequences defined in the specification, where the oligonucleotide comprises at least 15 nucleotides; or (b) a combination of an oligonucleotide comprising a sequence complementary to the complementary strand of a polynucleotide which comprises a 5'-end sequence and an oligonucleotide comprising a sequence complementary to a polynucleotide which comprises a 3'-end sequence, where the oligonucleotide comprises at least 15 nucleotides and the combination of oligonucleotide comprises at least 15 nucleotides and the combination of the 5'-end sequence/3'-end sequence is selected from those defined in the specification. The primer sets can be used in antisense therapy and in gene therapy. The primers are useful for synthesizing polynucleotides, particularly full-length cDNAs. The primers allow obtaining of the full-length detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs. The primers allow obtaining of the full-length cDNAs easily without any specialised methods. AAH03166 to AAH13628 and AAH13633 to AAH18742 represent human cDNA sequences; AAH92446 to AAH95893 represent human amino acid sequences; and AAH13629 to AAH13632 represent oligonucleotides, all of which are used in the exemplification of the present invention.
Sequence 1802 BP; 644 A; 298 C; 395 G; 465 T; 0 other;

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844 GGGACTGGTCCAGAGAGGCTGCCGTTTAACTTTTATCTCGTGAGAAGT 893
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944 CCACGAGCTTCAATACTTTAAGATCATTTATACAAAGACAAAGAAAGGT 993
67 lAlaIleValGluGluLeuValValGlyTyrGluThrSerLeuLysSerC 84
994 GGCAATCATAGAACAGCTAGTAGTAGGTATGAAACCTCTCTAAAGAGCT 1043
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1794 CCCAGATA 1801
  
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XX AC AAH16424;

XX DT 26-JUN-2001 (first entry)

XX DE Human cDNA sequence SEQ ID NO:15407.

XX KW Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.

XX OS Homo sapiens.

XX PN EP1074617-A2.

XX PD 07-FEB-2001.

XX PF 28-JUL-2000; 2000EP-0116126.

XX PR 29-JUL-1999; 95JP-0248036.

XX PR 27-AUG-1999; 95JP-0300253.

XX PR 11-JAN-2000; 2000JP-0118776.

XX PR 02-MAY-2000; 2000JP-0183767.

XX PR 09-JUN-2000; 2000JP-0241899.

XX PA (HELI-) HELIX RES INST.

XX PI Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;

XX PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;

XX DR WPI; 2001-318749/34.

XX PT Primer sets for synthesizing polynucleotides, particularly the 5602

XX PT full-length cDNAs defined in the specification, and for the detection

XX PT full-length cDNAs -

XX PS Claim 8; SEQ ID 15407; 2537pp + CD ROM; English.

XX CC The present invention describes primer sets for synthesizing 5602

XX CC full-length cDNAs defined in the specification. Where a primer set

XX CC comprises: (a) an oligo-dr primer and an oligonucleotide complementary

XX CC to the complementary strand of a polynucleotide which comprises one of

XX CC the 5602 nucleotide sequences defined in the specification, where the

XX CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination

XX CC of an oligonucleotide comprising a sequence complementary to the

XX CC complementary strand of a polynucleotide which comprises a 5'-end

XX CC sequence and an oligonucleotide comprising a sequence complementary to a

XX CC polynucleotide which comprises a 3'-end sequence, where the

CC oligonucleotide comprises at least 15 nucleotides and the combination of
 CC the 5'-end sequence/3'-end sequence is selected from those defined in
 CC the specification. The primer sets can be used in antisense therapy and
 CC in gene therapy. The primers are useful for synthesizing polynucleotides,
 CC particularly full-length cDNAs. The primers are also useful for the
 CC detection and/or diagnosis of the abnormality of the proteins encoded by
 CC the full-length cDNAs. The primers allow obtaining of the full-length
 CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
 CC AAH13633 to AAH18742 represent human cDNA sequences; AAH92446 to
 CC AAH95893 represent human amino acid sequences; and AAH13629 to AAH13632
 CC represent oligonucleotides, all of which are used in the exemplification
 CC of the present invention.

XX Sequence 1985 BP: 652 A; 349 C; 492 G; 492 T; 0 other;

alignment_scores: Quality: 1521.50 Length: 298
 Ratio: 5.175 Gaps: 1
 Percent Similarity: 98.658 Percent Identity: 97.315

alignment_block:

US-09-836-410A-1 x AAH16424 ..

Align seg 1/1 to: AAH16424 from: 1 to: 1985

1 MetLeuGluArgLeuLysIleTyrGluAlaThrThrLysTyrProAr 17
 1096 ATGTTAGAACGGCTAAAAAATTTATGAGGAAGCGCTGGACTTAATATCCAG 1145
 17 gGlyLeuValProArgLysLeuProLeuAsnPheLeuSerGlyGlyLysP 34
 1146 GGGACTGGTGCCCAAGAGCGCTGCGCTTAACCTTTTATCTGGTGAGAAGT 1195
 34 heLysGluCysLeuAspArgPheLeuArgMetAsnPheSerLysGlyCys 50
 1196 TTAAAGATGTTTGGATAGTTTCTTCAAGATGAATTTTCAGCAAGGGTTGC 1245
 51 ProProValPheAsnThrLeuAspSerLeuTyrArgAspLysGluLysVa 67
 1246 CCACCAGCTTCAATACCTTTTATGATCATTTATCAAGACAAAGAAAGGT 1295
 67 lAlaIleValGluGluLeuValValGlyTyrGluThrSerLeuLysSerC 84
 1296 GCATCATAGAGAGGTAGTAGTAGTATGAGTATGAAACCTCTCTAAAGAGCT 1345
 84 ysArgLeuPheAsnProAsnAspAspGlyLysGluGluProProThrThr 100
 1346 GCGGCTTATTTAACCCCAATGATGATGGAAGGAGGAGGACCCCAACACCA 1395
 101 LeuLeuTrpValGlnTyrTyrLeuAlaGlnHisTyrAspLysIleGlyG 117
 1396 TTACTTTGGGTCCAGTACTTCTGGCACAAACATTATGACAAAATTGGTCA 1445
 117 nProSerIleAlaLeuGluTyrIleAsnThrAlaIleGluSerThrProT 134
 1446 GCCATCTATTGCTTTGGAGTACATAAATCTGCTATTGAAAGTACACCTA 1495
 134 hrLeuIleGluLeuPheLeuValLysAlaLysIleTyrLysHisAlaGly 150
 1496 CATTAATAGAACCTTTCTCGTGAAGAGCTTAAATCTATAGCATGCTGGA 1545
 151 AsnIleLysGluAlaAlaArgTrpMetAspGluAlaGlnAlaLeuAspTh 167
 1546 AATATTAAAGAGCTCAAGGTGGATGGATGAGGCCAGGCCCTTGGACAC 1595
 167 rAlaAspArgPheIleAsnSerLysCysAlaLysTyrMetLeuLysAla 184
 1596 AGCAGACAGATTTATCAACTCCAAATGTGCAAAATACATCTTAAAGCCCA 1645
 184 snLeuIleLysGluAlaGluMetCysSerLysPheThrArgGlyGly 200
 1646 ACCTGATTAAAGAGCTGAGAAATGTGCTCAAGTTTACAGGGAAGGA 1695

201 ThrSerAlaValGluAsnLeuAsnGlnMetGlnCysMetTrpPheGlnTh 217
 1696 ACATCAGCGGTAGAGAATTTGAATGAAATGAGTGCATGTGGTTCCAAAC 1745
 217 tGluCysAlaGlnAlaTyrLysAlaMetAsnLysPheGlyGluAlaLeuL 234
 1746 AGAATGTGCCAGGCTTAAAGCAATGAATAAATTTGGTGAAGCACTTA 1795
 234 yLysCysHisGlnIleGluArgHisPheIleGluIleThrAspAspGln 250
 1796 AGAATGTGCATGAGATTGAGAGAAA.....ATCACTGATGACCAG 1835
 251 PheAspPheHisThrTyrCysMetArgLysIleThrLeuArgSerTyrVa 267
 1836 TTTGACTTTTCATACATACATGCTATGAGGAAGATTACCTTAGATCATATG 1885
 267 lAspLeuLeuLysLeuGluAspValLeuArgGlnHisPropheTyrPheL 284
 1886 GGACTTATTAAACTAGAGATGCTACTTCGACAGCATCCATTTACTTCA 1935
 284 ysAlaAlaArgIleAlaIleGluIleTyrLeuLysLeuHisasp 298
 1936 AGGCAGCAAGATTGCTATAGAGATCTATTGAAGCTTCATGAC 1979

seq_name: /SIDS1/gcgdata/hold-geneseq/geneseqn-emb1/NA2001A.DAT:AAH1664

seq_documentation_block:
 ID AAH1664 standard: DNA; 710 BP.

XX AAH1664;

XX 21-SEP-2001 (first entry)

XX Human differential transcription-associated cDNA SEQ ID 173.

XX Differential transcription; human; rat; tumour cell; cytostatic;
 KW Ras modulator; Class II tumour suppressor gene; gene therapy; ss.

XX Homo.sapiens.

XX WO200157058-A2.

XX 09-AUG-2001.

XX 31-JAN-2001; 2001WO-EP01003.

XX 31-JAN-2000; 2000DE-1004102
 XX (META-) METAGEN GES GENOMFORSCHUNG MBH.

XX Rosenthal A, Hinzmann B, Schaefer R, Zuber J, Tchernitsa O;
 PI Grips M, Hellriegel M, Schmitz A, Sers C;

XX WPI; 2001-483415/52.

XX Nucleic acids differentially expressed between tumor and normal cells,
 PT useful for diagnosis or therapy of tumors and for screening active
 PT agents

XX Disclosure: Page 376; 579pp; German.

XX This invention describes a nucleic acid (I) with differential expression
 CC between tumour and normal cells and which has cytostatic activity. (I)
 CC work as modulators of Ras activity by inducing expression of tumour
 CC suppressor genes. (I), and polypeptides encoded by them, are useful as
 CC targets for diagnosis or therapy and in screening to determine the
 CC effects of an active compound (potential pharmaceutical) on a cell line,
 CC particularly for diagnosis and treatment of tumors, especially by
 CC modulating expression of (I) (by gene therapy, antisense RNA or ribozyme
 CC methods) or by modulating the amount and/or location of (I)-encoded
 CC polypeptides (by administration of the polypeptide or its activator,
 CC antibody (optionally as a conjugate) or inhibitor). The method allows

CC Identification of many Class II tumour suppressor genes (i.e. genes that
 CC are not primary targets for tumour-initiating mutations).
 CC AAH81492-AAH82376 represent the human and rat derived nucleic acid
 CC fragments described in the method of the invention.
 XX
 SQ Sequence 710 BP; 170 A; 146 C; 121 G; 267 T; 6 other;

alignment_scores:
 Quality: 1176.00 Length: 237
 Ratio: 5.091 Gaps: 0
 Percent Similarity: 97.468 Percent Identity: 96.203

alignment_block:

US-09-836-410A-1 x AAH81664/rev ..

Align seg 1/1 to reverse of: AAH81664 from: 1 to: 710

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103 TrpValGlnTyrTyrLeuAlaGlnHisTyrAspLysIleGlyGlnProse 119
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710 TGGGTCAAGTACTTGGCACACCATTTATGACAAAATTGGTCAGCCATC 661
119 rIleAlaLeuGluTyrIleAsnThrAlaIleGluSerThrProThrLeuI 136
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
660 TANTGCTNTGGAGTACATAAATCTGCTATTGANAGTACACT ACATTA 612
136 leGluLeuPheLeuValLysAlaLysIleTyrLysHisAlaGlyAsnIle 152
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
611 TAGACNTCTTCTCGTGANAGCTAAATCTATAAGCATGCTGGAATAAT 562
153 LysGluAlaAlaArgTrpMetAspGluAlaGlnAlaLeuAspThrAlaAs 169
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
561 AAAGAAGCTGCAAGTGGATGGATGAGGCGCCCTTGGACACACGACA 512
169 pArgPheIleAsnSerLysCysAlaLysTyrMetLeuLysAlaAsnLeuI 186
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
511 CAGATTATCAACTCCAAATGTGCAAAATACATGCTAAAGGCAACCTGA 462
186 leLysGluAlaGluMetCysSerLysPheThrArgGluGlyThrSer 202
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
461 TTAAGAAGCTGAAGAAATGTCTCAAGTTTACAGGGAAGCAACATCA 412
203 AlaValGluAsnLeuAsnGluMetGlnCysMetTrpPheGlnThrGluCy 219
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
411 GCGGTAGAGAAATTTGAATGAATGCAATGTCATGTGTTCCAAACAGATG 362
219 sAlaGlnAlaTyrLysAlaMetAsnLysPheGlyGluAlaLeuLysLysC 236
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
361 TGCCCGAGGCTTATAAGCAATGAATAATTTGTTGAAGCACCTTAAGAA 312
236 yHisGluIleGluArgHisPheIleGluIleThrAspAspGlnPheAsp 252
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
311 GTCATGAGATTGAGAGACATTTTATAGAAATCACTGATGACCATTTGAC 262
253 PheHisThrTyrCysMetArgLysIleThrLeuArgSerTyrValAspLe 269
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
261 TTTTACATACATCTGTAGGAGAGATTACCCCTTAGATCATATGTGACTT 212
269 uLeuLysLeuGluAspValLeuArgGlnHisProPheTyrPheLysAlaA 286
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
211 ATTAAGTACAGAGATGCTTTCGACACATCCATTTTACTTCAAGGAG 162
286 laArgIleAlaIleGluIleTyrLeuLysLeuHisAspAsnProLeuThr 302
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
161 CAAGAATTGCTATAGATCATCTTTTGAAGCTTCATGACAAACCCCTTACA 112
303 AspGluAsnLysGluHisGluAlaAspThrAlaAsnMetSerAspLysG 319
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
111 CATGAGAATAAAGAACAGGAGCTGATACAGCAACATGTCTGACAAAGA 62
319 uLeuLysLysLeuArgAsnLysGlnArgAlaGlnLysLysAlaGlnI 336
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
61 GCTAAGAAGCTACGTATATAACAAAGAGAGCTCAAAAGAGCCCCAGA 12

```

336 leGluGluGlu 339
 |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
 11 TAGAAGAAGAG 1

seq_name: /SIDSI/gcdata/hold-geneseq/geneseqn-emb1/NA1999.DAT:AAZ15705

seq_documentation_block:

XX ID AAZ15705 standard; cDNA; 781 BP.

XX AC AAZ15705;

XX DT 12-OCT-1999 (first entry)

XX DE Human gene expression product cDNA sequence SEQ ID NO:3174.

XX KW Human; gene: gene expression product; diagnosis; therapy; probe;
 KW detection; mapping; tissue typing; profiling; forensic; cancer;
 KW genetic analysis; colorectal cancer; breast cancer; lung cancer; ss.
 XX OS Homo sapiens.

XX PN WO9938972-A2.

XX PD 05-AUG-1999.

XX XX 28-JAN-1999; 99WO-US01619.

XX PF 03-APR-1998; 98US-0080666.

XX PR 28-JAN-1998; 98US-0072910.

XX PR 24-FEB-1998; 98US-0075954.

XX PR 31-MAR-1998; 98US-0080114.

XX PR 03-APR-1998; 98US-0080515.

XX PA (CHIR) CHIRON CORP.

XX PA (HYSE-) HYSEQ INC.

XX PI Crkvenjakov R, Dickson M, Drmanac R, Drmanac S;
 PI Escobedo J, Garcia PD, Garcia V, Glese K, Innis MA;
 PI Jones WL, Kassam A, Kennedy GC, Kita D, Labat I;
 PI Lamson G, Leshkowitz D, Pot D, Randazzo F, Reinhard C;
 PI Stache-Crain B, Sudduth-Klinger J, Williams LT;

XX DR WPI; 1999-494092/41.

XX PT Novel human genes and their expression products which are
 PT differentially expressed in different cell types

XX XX Claim 1; Page 1524-1525; 2479pp; English.

XX CC The present invention describes a library of human polynucleotides
 CC comprising the sequences given in AAZ1532 to AAZ1779. Also described is
 CC a method of detecting differentially expressed genes correlated with the
 CC cancerous state of a mammalian cell, comprising detecting at least one
 CC differentially expressed gene product in a test sample from a cell
 CC suspected of being cancerous, where the gene product is encoded by one
 CC of the 5248 polynucleotide sequences given in AAZ1532 to AAZ1779. The
 CC polynucleotides can be used as a source of primers and probes, which can
 CC be used for a variety of purpose, e.g. detection of expression levels,
 CC mapping, tissue typing or profiling, forensics, genetic analysis and
 CC detection of polymorphisms. Polypeptides encoded by the polynucleotides
 CC can be used for raising antibodies for experimental, diagnostic and
 CC therapeutic purposes. The polynucleotides may also be used to construct
 CC arrays for diagnostics (which may be used to determine function of an
 CC encoded protein); and to detect differences in expression levels between
 CC two cells (e.g. to identify abnormal or diseased tissue in a human, to
 CC identify a genetic predisposition or susceptibility to a disease such as
 CC cancer). The polynucleotides of the invention are especially used in the
 CC diagnosis, prognosis and management of colorectal cancer, breast cancer,
 CC and lung cancer. The polynucleotides can also be used to screen for
 CC peptide analogues and antagonists.

XX SQ Sequence 781 BP; 263 A; 140 C; 156 G; 205 T; 17 other;

```

alignment_scores:
  Quality: 1104.00      Length: 232
  Ratio: 4.973          Gaps: 4
  Percent Similarity: 95.690  Percent Identity: 95.259

alignment_block:
  US-09-836-410a-1 x AAZ15705 ..

Align seg 1/1 to: AAZ15705 from: 1 to: 781

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|||||
87  AGAGACAAGAAAGAGTGGCAATCATAGAGAGTGTAGTAGGTATG 136
|||||

78  luThrSerLeuLysSerCysArgLeuPheAsnProAsnAspGlyLys 94
|||||
137  AAACCTCTCTAAAAGCTGCCGTTATTTAAACCCCAATGATGGAAG 186
|||||

95  GluGluProProThrThrLeuLeuTrpValGlnTyrTyrLeuAlaGlnHi 111
|||||
187  GAGAACCAACCAACCAATCTACTTTGGTCCNNTNCTACTTGGCAACA 236
|||||

111  sTyrAspLysIleGlnProSerIleAlaLeuGluTyrIleAsnThrA 128
|||||
237  TTATGACAAAATTTGGTCAGCCATCTATTGCTTGGAGTACATAAATCTG 286
|||||

128  laIleGluSerThrProThrLeuLeuGluLeuPheLeuValLysAlaLys 144
|||||
287  CTATTGAAAGTACACCTACATTAATAGAACTCTTTCTCGTGAAGCTTAA 336
|||||

145  IleTyrLysHisAlaGlyAsnIleLysGluAlaAlaArgTrpMetAspG1 161
|||||
337  ATCTATAGCATGCTGGAATATTAAAGAGCTGCAGGTGGATGATGA 386
|||||

161  uAlaGlnAlaLeuAspThrAlaAspArgPheIleAsnSerLysCysAlaL 178
|||||
387  GGCCAGGCGCTTGGACAGCAGACAGATTTATCAACTCCAAATGTGCAA 436
|||||

178  ysTyrMetLeuLysAlaAsnLeuIleLysGluAlaGluMetCysSer 194
|||||
437  AATACATGCTAAAGCCCAACCTGATTAAAGAGCTGAAGAAATGTGCTCA 486
|||||

195  LysPheThrArgGluGlyThrSerAlaValGluAsnLeuAsnGluMetG1 211
|||||
487  AAGTTTACAGGGAAGGACATCAGCGTAGAGATTTGAATGAATGCA 536
|||||

211  nCysMetTrpPheGlnThrGluCysAlaGlnAlaTyrLysAlaMetAsnL 228
|||||
537  GTGCATGTGGTTCCAAACAGAAATGTGCCAGGCTTATAAAGCAATGAATA 586
|||||

228  ysPheGlyGluAlaLeuLysCysHisGluLeGluArgHisPheIle 244
|||||
587  AATTTGGTGAAGCACTTAAGAAATGTCATGAGATTCAGAGACATTTTATA 636
|||||

245  GluIleThrAspAspGlnPheAspPheHisThrTyrCysMet.ArgLys 260
|||||
637  GGAATCATCTGATGACCATTTGACTTTTCATACATCTGATGGAAGGAAG 686
|||||

261  IleThrLeuArgSerTyrValAspLeuLeuLysLeu.GluAspValLeuA 277
|||||
687  ATTACCTTAGATCATATGTGGACTTATTNAACTATGAAGAATGACTTT 736
|||||

277  rGlnHisProPheTyrPheLysAlaAlaArgIleAla 289
|||||
737  NCAGCATNCATTTTACTTTCAGGCGCAGCAAGATTGCT 774
|||||

seq_name: /SIDS1/gcgdata/hold-geneseq/geneseqn-emb1/NAL1999.DAT:AAZ99053
seq_documentation_block:
ID  AAZ99053 standard; cDNA: 781 BP.
XX

```

```

AC  AAX99053;
XX
XX  24-SEP-1999 (first entry)
DE  Human validated cancer cell derived cDNA #375.
XX
XX  Cancer; human; colon; breast; lung; transmembrane receptor; ATPase;
KW  integral membrane protein; aspartyl protease; GATA family; wnt family;
KW  transcription factor; G-protein alpha subunit; protein phosphatase;
KW  phospholipase; tyrosine phosphatase; diacylglycerol binding protein; trypsin;
KW  protein kinase; tyrosine phosphatase; developmental signalling protein;
KW  WW/rsp5/WWP domain; therapy; forensic; genetic mapping; diagnostic;
KW  detection; treatment; cervical; melanoma; colorectal adenocarcinoma;
KW  Wilm's tumour; retinoblastoma; sarcoma; myosarcoma; lung carcinoma;
KW  leukemia; lymphoma; dysplasia; hyperplasia; endometrium; adrenal;
KW  prostate; ss.
XX
XX  Homo sapiens.
XX
XX  WO9933982-A2.
PN  08-JUL-1999.
XX
XX  22-DEC-1998; 98WO-US27610.
XX
XX  21-DEC-1998; 98US-0217471.
PR  23-DEC-1997; 97US-0058755.
PR  03-APR-1998; 98US-0080664.
PR  21-OCT-1998; 98US-0105234.
PR  27-OCT-1998; 98US-0105877.
XX
XX  (CHIR ) CHIRON CORP.
PA  (HYSE-) HYSEQ INC.
XX
XX  CrkVenjakov R, Dickson M, Drmanac R, Drmanac S;
PI  Escobedo J, Garcia PD, Garcia V, Giese K, Innis M;
PI  Jones LW, Kassam A, Kennedy GC, Kita D, Labat I;
PI  Lamson G, Leshkowitz D, Pot D, Randazzo F, Reinhard C;
PI  Stache-Crain B, Sudduth-Klinger J, Williams LT;
XX
XX  WPI: 1999-430243/36.
XX
XX  New isolated human polynucleotides
XX
XX  Claim 1: Page 564; 591pp; English.
XX
XX  This invention describes novel isolated human polynucleotides obtained
CC  by screening for differential expression in colon cancer, breast cancer
CC  and lung cancer cell lines. The polynucleotides of the invention are
CC  represented in AAX98275-X99118 and encode polypeptides of protein
CC  families selected from 4 transmembrane segments integrated with various
CC  proteins, 7 transmembrane receptors, ATPases associated with various
CC  cellular activities (AAA), eukaryotic aspartyl proteases, GATA family of
CC  transcription factors, G-protein alpha subunit, phospholipase or
CC  diacylglycerol binding proteins, protein kinase, protein phosphatase 2C,
CC  protein tyrosine phosphatase, trypsin, wnt family of developmental
CC  signalling proteins and WW/rsp5/WWP domain containing proteins. The
CC  encoded polypeptides also have a functional domain selected from Ank
CC  repeat, basic region plus leucine zipper transcription factors,
CC  bromodomain, EF-hand, SH3 domain, WD domain/G-beta repeats, zinc finger
CC  (C2H2 type), zinc finger (CCHC class), and zinc-binding metalloprotease
CC  domain. The polynucleotides encode polypeptides with similarity to known
CC  protein families and are predicted to have similar properties. The novel
CC  polynucleotides can be used to develop products for use as therapeutic
CC  agents and in forensics, genetic analysis, mapping and diagnostic
CC  applications. In particular the product can be used for the detection
CC  and management of cancers. They can be used for treating e.g. cervical
CC  cancers, melanomas, colorectal adenocarcinomas, Wilm's tumour, sarcomas,
CC  retinoblastoma, myosarcomas, lung carcinomas, monocytic leukemia, and
CC  myelogenous leukemia, promyelocytic leukemia, histiocytic lymphoma, anhydric
CC  myeloid leukemia, and lymphomas such as histiocytic lymphoma, anhydric
CC  hereditary ectodermal dysplasia, congenital alveolar dysplasia, and
CC  epithelial dysplasia of the cervix, fibrous dysplasia of bone, and

```

CC mammary dysplasia, hyperplasias, e.g. endometrial, adrenal, breast,
CC prostate or thyroid hyperplasias or pseudoeplitheliomatous hyperplasia of
CC the skin.
XX
SQ Sequence 781 BP; 263 A; 140 C; 156 G; 205 T; 17 other;

alignment_scores:
Quality: 1104.00 Length: 232
Ratio: 4.973 Gaps: 4
Percent Similarity: 95.690 Percent Identity: 95.259

alignment_block:

US-09-836-410A-1 x AAX99053 ..

Align seg 1/1 to: AAX99053 from: 1 to: 781

62 ArgAspLysGlu.LysValAlaIleValGluLeuValValcIlyTyrG 78
87 AGAGACAAGAAAAGGTGGCAATCATAGAAGAGTTAGTAGGTATG 136
78 luThrSerLeuLysSerCysArgLeuPheAsnProAsnAspGlyLys 94
137 AACCTCTCTAAAAGCTGCGGTATTTAAACCCCAATGATGAAAG 186
95 GluGluProProThrThrLeuLeuTrpValGlnTyrTyrLeuAlaGlnHi 111
187 GAGAACCAACCAACCATACATTTGGTCCNNTACTTGGCCACAACA 236
111 sTyrAspLysIleGlyGlnProSerIleAlaLeuGluTyrIleAsnThra 128
237 TTATGACAAAATTTGGTCAGCCATCTATTGCTTTGGAGTACATAAATCTG 286
128 laileGluSerThrProThrLeuIleGluLeuPheLeuValLysAlaLys 144
287 CTATTGAAGTACACCTACATTAATAGAACTCTTCTCGTGAAGCTAAA 336
145 IletTyrLysHisIaGlyAsnIleLysGluAlaAlaArgTrpMetAspGI 161
337 ATCTAATGACATGCTGGAATATTAAGAAGCTGCAAGTGGATGATCA 386
161 uAlaGlnAlaLeuAspThrAlaAspArgPheIleAsnSerLysCysAlaL 178
387 GGCCAGGCTTGGACACAGCAGACAGATTTATCAACTCCAAATGTGCAA 436
178 ySTyrMetLysAlaAsnLeuIleLysGluAlaGluGluMetCysSer 194
437 AATACATGCTAAAGGCAACCTGATTAAAGAGCTGAAGAAATGTCTCA 486
195 LysPheThrArgGluGlyThrSerAlaValGluAsnLeuAsnGluMetGI 211
487 AAGTTTACAGGGAAGGAACATCAGCGGTAGAGAATTTGAATGAATGCA 536
211 nCysMetTrpPheGlnThrGluCysAlaGlnAlaTyrLysAlaMetAsnL 228
537 GTGCATGTGTTCCAAACAGAAATGTCCCGAGCTTATAAGCAATGAATA 586
228 yPheGlyGluAlaLeuLysCysHisGluIleGluArgHisPheIle 244
587 AATTTGGTGAAGCACTTAAGAAATGTCATGAGATGAGAGACATTTTATA 636
245 GluIleThrAspAspGlnPheAspPheHisThrTyrCysMet.ArgLys 260
637 GGAATCACTGATGACCACTTTGACTTTTCATACATCTGATGAGGAAG 686
261 IleThrLeuArgSerTyrValAspLeuLeuLysLeu.GluAspValLeuA 277
687 ATTACCCCTTAGATCATATGTGGACTTATTNAACTATGAGATGTACTTT 736
277 rGlnHisProPheTyrPheLysAlaAlaArgIleAla 289
737 NACAGCATNCATTTTACTTCAAGCAGCAAGAATGTCT 774

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seq_documentation_block:
ID AAZ15983 standard; CDNA: 764 BP.
XX
AC AAZ15983;
XX
DT 12-OCT-1999 (first entry)
XX
DE Human gene expression product cDNA sequence SEQ ID NO:3452.
XX
KW Human; gene; gene expression product; diagnosis; therapy; probe;
KW detection; mapping; tissue typing; profiling; forensic; cancer;
KW genetic analysis; colorectal cancer; breast cancer; lung cancer; ss.
XX
OS Homo sapiens.
XX
XX WO9938972-A2.
XX 05-AUG-1999.
XX
PD 28-JAN-1999; 99WO-US01619.
XX
PF 03-APR-1998; 98US-0080666.
XX
PR 28-JAN-1998; 98US-0072910.
XX
PR 24-FEB-1998; 98US-0075954.
XX
PR 31-MAR-1998; 98US-0080114.
XX
PR 03-APR-1998; 98US-0080515.
XX
PA (CHIR) CHIRON CORP.
XX
PA (HYSE-) HYSEQ INC.
XX

PI Crkvenjakov R, Dickson M, Drmanac R, Drmanac S;
PI Escobedo J, Garcia PD, Garcia V, Giese K, Innis MA;
PI Jones WL, Kassam A, Kennedy GC, Kita D, Labat J;
PI Lamson G, Leshkowitz D, Pot D, Randazzo F, Reinhard C;
PI Stache-Crain B, Sudduth-Klinger J, Williams LT;
XX
DR WFI: 1999-494092/41.
XX
Novel human genes and their expression products which are
differentially expressed in different cell types
Claim 1: Page 1650-1651; 2479pp; English.
XX

The present invention describes a library of human polynucleotides
comprising the sequences given in AAZ12532 to AAZ1779. Also described is
a method of detecting differentially expressed genes correlated with the
cancerous state of a mammalian cell, comprising detecting at least one
differentially expressed gene product in a test sample from a cell
suspected of being cancerous, where the gene product is encoded by one
of the 5248 polynucleotide sequences given in AAZ12532 to AAZ1779. The
polynucleotides can be used as a source of primers and probes, which can
be used for a variety of purpose, e.g. detection of expression levels,
mapping, tissue typing or profiling, forensics, genetic analysis and
detection of polymorphisms. Polypeptides encoded by the polynucleotides
can be used for raising antibodies for experimental, diagnostic and
therapeutic purposes. The polynucleotides may also be used to construct
arrays for diagnostics (which may be used to determine function of an
encoded protein); and to detect differences in expression levels between
two cells (e.g. to identify abnormal or diseased tissue in a human, to
identify a genetic predisposition or susceptibility to a disease such as
cancer). The polynucleotides of the invention are especially used in the
diagnosis, prognosis and management of colorectal cancer, breast cancer,
and lung cancer. The polynucleotides can also be used to screen for
peptide analogues and antagonists.

Sequence 764 BP; 253 A; 138 C; 148 G; 204 T; 21 other;

alignment_scores:
Quality: 973.00 Length: 200
Ratio: 5.015 Gaps: 3

Percent similarity: 97.000 Percent Identity: 95.500

alignment_block:

US-09-836-410A-1 x AAZ15983

Align seg 1/1 to: AAZ15983 from: 1 to: 764

62 ArgAspLysGluLysValAlaIleValGluLeuValValGlyTyrG1 78
|||||
65 AGACAAAGAAAGGCGCAATCATAGAGAGTTTACTAGGTTATGA 114
|||||
78 uThrSerLeuLysSerCysArgLeuPheAsnProAsnAspGlyLysG 95
|||||
115 AACCTCTCTAAAGAGCTGCCGGTTATTTAACCCCAATGATGGAAGG 164
|||||
95 LuGluProThrThrLeuLeuTrpValGlnTyrTyrLeuAlaGlnHis 111
|||||
165 AGGAACCAACCAACCATTAATTTGGGTCCAGTACTACTTGGCACAACAT 214
|||||
112 TyrAspLysIleGlyGlnProSerIleAlaLeuGluTyrIleAsnThrAl 128
|||||
215 TATGACAAATTTGGTCAGCCATCTATTGCTTTGGAGTACATAATCTGC 264
|||||
128 aIleGluSerThrProThrLeuIleGluLeuPheLeuValLysAlaLysI 145
|||||
265 TATTGAAAGTACACTACATTAATAGAACTCTTCTCGTGAAGAGCTTAAA 314
|||||
145 LeTyrLysHisAlaGlyAsnIleLysGluAlaAlaArgTrpMetAspGlu 161
|||||
315 TCTATAAGCATGCTGGAATATTAAAGAGCTGCAAGGTGGATGATGAG 364
|||||
162 AlaGlnAlaLeuAspThrAlaAspArgPheIleAsnSerLysCysAlaLy 178
|||||
365 GCCCAGGCTTGGACACAGCAGACAGATTTATCACTCCCAATGTGCNAA 414
|||||
178 sTyrMetLeuLysAlaAsnLeuIleLysGluAlaGluMetCysSerL 195
|||||
415 ATACATGTCTAAAGCAACCTGATTAAGAGAGCTGAAGAAATGTCTCAA 464
|||||
195 yspPheThrArgGluThrSerAlaValGluAsnLeuAsnGluMetGln 211
|||||
465 AGTTTACAGGGAAGGAACATCAGCGGTAGAGAAATTTGAAATGAATGAG 514
|||||
212 CysMetTrpPheGluThrGluCysAlaGlnAlaTyrLysAlaMetAsn.L 228
|||||
515 TGCATGTGTTTCCAAACAGAAATGTCGCCAGGCTTATAAGCAATGAATA 564
|||||
228 yspPheGluAlaLeuLysLysCysHis.GluIleGluArgHisPheIl 244
|||||
565 AATTTGGTGAAGCACTTAAGAAATGTCATTTGAGATTGAGAGACTTTTATA 614
|||||
244 eGluIleThrAspAsp.GlnPheAspPheHisThrTyrCysMet 258
|||||
615 GGAAATTCATGATGACCCAGTTGACTTTTCATACATACATCTGATG 658
|||||

seq_name: /SIDS1/gcgdata/hold-geneseq/geneseqn-emb1/NA1999.DAT:AA98777

seq_documentation_block:

ID AA98777 standard; cDNA; 764 BP.

XX

AC

XX

XX

DE

XX

XX

XX

XX

Cancer; human; colon; breast; lung; transmembrane receptor; ATPase; integral membrane protein; aspartyl protease; GATA family; wnt family; transcription factor; G-protein alpha subunit; protein phosphatase; phorbol ester binding protein; diacylglycerol binding protein; trypsin; protein kinase; tyrosine phosphatase; developmental signalling protein; WW/rsp5/WWP domain; therapy; forensic; genetic mapping; diagnostic; detection; treatment; cervical; melanoma; colorectal adenocarcinoma;

Wilm's tumour; retinoblastoma; sarcoma; myosarcoma; lung carcinoma; leukemia; lymphoma; dysplasia; hyperplasia; endometrium; adrenal; prostate; ss.

XX Homo.sapiens.

XX W09933982-A2.

XX 08-JUL-1999.

XX 22-DEC-1998; 98WO-US27610.

XX 21-DEC-1998; 98US-0217471.

XX 23-DEC-1997; 97US-0068755.

XX 03-APR-1998; 98US-0080664.

XX 21-OCT-1998; 98US-0105234.

XX 27-OCT-1998; 98US-0105877.

XX (CHIR) CHIRON CORP.

XX (HYSE-) HYSEQ INC.

XX Crkvenjakov R, Dickson M, Drmanac R, Drmanac S;

XX Escobedo J, Garcia PD, Garcia V, Giese K, Innis MA;

XX Jones LW, Kassam A, Kennedy GC, Kita D, Labat I;

XX Lamson G, Leshkowitz D, Pot D, Randazzo F, Reinhard C;

XX Stache-Crain B, Sudduth-Klinger J, Williams LT;

XX WPI; 1999-430243/36.

XX New isolated human polynucleotides

XX Claim 1: Page 453; 591pp: English.

This invention describes novel isolated human polynucleotides obtained by screening for differential expression in colon cancer, breast cancer and lung cancer cell lines. The polynucleotides of the invention are represented in AAX98275-X99118 and encode polypeptides of protein families selected from 4 transmembrane segments integral membrane proteins, 7 transmembrane receptors, ATPases associated with various cellular activities (AAA), eukaryotic aspartyl proteases, GATA family of transcription factors, G-protein alpha subunit, phorbol esters or diacylglycerol binding proteins, protein kinase, protein phosphatase 2C, protein tyrosine phosphatase, trypsin, wnt family of developmental signalling proteins and WW/rsp5/WWP domain containing proteins. The encoded polypeptides also have a functional domain selected from Ank repeat, basic region plus leucine zipper transcription factors, bromodomain, EF-hand, SH3 domain, WD domain/G-beta repeats, zinc finger (C2H2 type), zinc finger (CCHC class), and zinc-binding metalloprotease domain. The polynucleotides encode polypeptides with similarity to known protein families and are predicted to have similar properties. The novel polynucleotides can be used to develop products for use as therapeutic agents and in forensics, genetic analysis, mapping and diagnostic applications. In particular, the product can be used for the detection and management of cancers. They can be used for treating e.g. cervical cancers, melanomas, colorectal adenocarcinomas, Wilm's tumour, sarcomas, retinoblastoma, myosarcomas, lung carcinomas, monocytic leukemia, and myelogenous leukemia, and lymphomas such as histiocytic lymphoma, anhydric myeloid leukemia, and congenital alveolar dysplasia, hereditary ectodermal dysplasia, fibrous dysplasia of bone, and epithelial dysplasia of the cervix, fibrous dysplasia of breast, mammary dysplasia, hyperplasias, e.g. endometrial, adrenal, breast, prostate or thyroid hyperplasias or pseudoepitheliomatous hyperplasia of the skin.

XX Sequence 764 BP; 253 A; 138 C; 148 G; 204 T; 21 other;

alignment_scores:

Quality: 973.00 Length: 200

Ratio: 5.015 Gaps: 3

Percent Similarity: 97.000 Percent Identity: 95.500

alignment_block:

US-09-836-410A-1 x AAX98777

Align seg 1/1 to: AAX98777 from: 1 to: 764

```
62 ArgAspLysGluLysValAlaIleValGluLeuValValGlyTyrGI 78
|||||
65 AGACACAAGAAAGGTGGCAATCATAGACAGTTTNTAGTAGCTATGA 114
|||||
78 uThrSerLeuLysSerCysArgLeuPheAsnProAsnAspAspGlyLysG 95
|||||
115 AACCTCTCTAAAAGCTGCCGGTTATTAAACCCCAATCATGATGGAAGG 164
|||||
95 luGluProThrThrLeuLeuTyrValGlnTyrTyrLeuAlaGlnHis 111
|||||
165 AGAACCAACCAACACATTAATTTGGGTCCAGTACTTGGGCACACAT 214
|||||
112 TyrAspLysIleGlyGlnProSerIleAlaLeuGluTyrIleAsnThrAl 128
|||||
215 TATGACAAAATTGGTCAGCCATCTATTGCTTGGAGTACATAAATCTGC 264
|||||
128 aileGluSerThrProThrLeuIleGluLeuPheLeuValLysAlaLysI 145
|||||
265 TATTGAAGTACACCTACATTAATAGAACTCTTTCTCGTGAAGCTAAAA 314
|||||
145 leTyrLysHisAlaGlyAsnIleLysGluAlaAlaArgTyrMetAspGlu 161
|||||
315 TCTATAAGCATGCTGGAAATATTAAAGAACCTGCAAGTGGATGGATGAG 364
|||||
162 AlaGlnAlaLeuAspThrAlaAspArgPheIleAsnSerLysCysAlaLy 178
|||||
365 GCCCAGGCCCTTGGACACAGACAGACAGATTTATCAACTCCAAATGTGCAA 414
|||||
178 sTyrMetLeuLysAlaAsnLeuIleLysGluAlaGluMetCysSerL 195
|||||
415 ATACATGCTAAAAGCAACCTGATTAAAGAGCTGAAGAATGTGCTCAA 464
|||||
195 ysPheThrArgGluGlyThrSerAlaValGluAsnLeuAsnGluMetGln 211
|||||
465 AGTTTACAAGGGAAGGAACATCAGCGGTAGAGAAATTTGAATGAATGCAG 514
|||||
212 CysMetTyrPheGlnThrGluCysAlaGlnAlaTyrLysAlaMetAsn.L 228
|||||
515 TGATGTGGTTCCAAACAGANATGTGCCAGGCTTATAAAGCAATGAATTA 564
|||||
228 ysPheGlyGluAlaLeuLysLysCysHis.GluIleGluArgHisPheIl 244
|||||
565 AATTGTGTGAAGCATTAAAGAAATGTCAATGAGATTGAGAGACTTTTATA 614
|||||
244 eGluIleThrAspAsp.GlnPheAspPheHisThrTyrCysMet 258
|||||
615 GGAATCACTGATGACCCAGTTTGACTTTTCATACATACATCTGATG 658
|||||
```

seq_name: /SIDS1/gcgdata/hold-geneseq/geneseq-emb1/NA2001A.DAT.AAS37350

seq_documentation_block:

ID AAS37350 standard; cDNA; 402 BP.

AC AAS37350;

DT 17-DEC-2001 (first entry)

DE Novel human diagnostic and therapeutic gene #408.

KW Human; cancer; breast; lung; colon; prostate; cytostatic; diagnostic; ss.

OS Homo sapiens.

PN WO2001/66753-A2.

PD 13-SEP-2001.

PF 09-MAR-2001; 2001WO-US07787.

XX

PR 09-MAR-2000; 2000US-0188609.

XX (CHIR) CHIRON CORP.

PA (HYSE-) HYSEQ INC.

XX Williams LT, Escobedo J, Innis MA, Garcia PD, Sudduth-Klinger J;

PI Reinhard C, Randazzo F, Kennedy GC, Pot D, Kassam A, Lamson G;

PI Drmanac R, Crkvenjakov R, Dickson M, Drmanac S, Labat I;

XX Leshkowitz D, Kita D, Garcia V, Jones WL, Stache-Crain B;

DR WPI; 2001-530177/58.

XX New polynucleotides and polypeptides, useful for diagnosis and

PT treatment of breast, lung and colon cancer -

XX Claim 1; Page 698-699; 1193pp; English.

XX The invention relates to new polynucleotides and polypeptides, useful for diagnosis and treatment of breast, lung and colon cancer. The sequences can be used in detecting differentially expressed genes correlated with a cancerous state of a mammalian cell, comprising detecting at least one differentially expressed gene product in a test sample derived from a cell suspected of being cancerous. They can also be used to inhibit tumour growth by modulating expression of a gene product. AAS36943-AAS39338 represent novel human diagnostic and therapeutic coding sequences of the invention.

XX Sequence 402 BP; 142 A; 76 C; 85 G; 99 T; 0 other;

alignment_scores:

Quality: 680.00 Length: 130

Ratio: 5.231 Gaps: 0

Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-836-410A-1 x AAS37350 ..

Align seg 1/1 to: AAS37350 from: 1 to: 402

71 GluGluLeuValValGlyTyrGluThrSerLeuLysSerCysArgLeuPh 87

|||||

12 GAAGAGTAGTAGTAGTTATGAACCTCTCTAAAGAGTCCCGGTATTT 61

|||||

87 eAsnProAsnAspAspGlyLysGluGluProProThrThrLeuLeuTyrPv 104

|||||

62 TAACCCCAATGATGATGGAAGAGGAGAACCAACCAACCACTTACTTTGGG 111

|||||

104 aGlnTyrTyrLeuAlaGlnHisTyrAspLysIleGlyGlnProSerIle 120

|||||

112 TCCAGTACTACTTGGCACACACATTTATGACAAATTTGGTCAGCCATCTATT 161

|||||

121 AlaLeuGluTyrIleAsnThrAlaIleGluSerThrProThrLeuIleGl 137

|||||

162 GCTTTGGAGTACATAAATACCTGCTATTGAAAGTACACCTACATTAATAGA 211

|||||

137 uLeuPheLeuValLysAlaLysIleTyrLysHisAlaGlyAsnIleLysG 154

|||||

212 ACTCTTCTCGTGAAGCTAAATCTATAAGCATGCTGGAATATTTAAAG 261

|||||

154 luAlaAlaArgTyrMetAspGluAlaGlnAlaLeuAspThrAlaAspArg 170

|||||

262 AAGCTGCAAGTGGATGGATGGCCAGGCTTTGGACACAGACAGACAGA 311

|||||

171 PheIleAsnSerLysCysAlaLysTyrMetLeuLysAlaAsnLeuIleLy 187

|||||

312 TTTATCAACTCCAAATGTGCAAAATACATGCTAAAGCCACCTGATTA 361

|||||

187 sGluAlaGluMetCysSerLysPheThrArgGluGly 200

|||||

362 AGAAGCTGAAGAAATGTGCTCAAAAGTTTACAAGGGAAGGA 401

|||||

seq_name: /SIDS1/gcgdata/hold-geneseq/geneseq-emb1/NA2001A.DAT.AAH12222

Align seq 1/1 to reverse of: AAH12222 from: 1 to: 488

142 LysAlaLysIleTyrLysHisAlaGlyAsnIleLysGluAlaAlaArgTr 158
466 GAAGCTAAATCTATAAGCA.CCTGNAANTTTNAAAGAGCCCTNCAAGGG 418
158 pMetAspGluAlaGlnAlaLeuAspThrAlaAspArgPheIleAsnSerL 175
417 GATGGATTAGGCCAGCCAGCCAGCCAGCCAGCCAGCCAGCCAGCCAGCC 368
175 yCysAlaLysTyrMetLeuLysAlaAsnLeuLysGluAlaGluGlu 191
367 AATGTGCAAAATACATGCTAAAGCCCAACCTGATTAAAGAACTGAAGA 318
192 MetCysSerLysPheThrArgGluGlyThrSerAlaValGluAsnLeuAs 208
317 ATGTGCTCAAGTTTNCAGGGAAGCAACATCAGCGGTANAGAAATTTGAA 268
208 nGluMetGlnCysMetTrpPheGlnThrGluCysAlaGlnAlaTyrLysA 225
267 TGAATGTCAGCATGNGGTTCCAAAAGCAATGNGCCAGGCTTATAAG 218
225 laMetAsnLysPheGlyGluAlaLeuLysCysHisGluIleGluArg 241
217 CAATGAATAAATTTGGTGAAGCNCCTTAAGAAATGTCATGAGATTGAGA 168
242 HisPheIleGluIleThrAspAspGlnPheAspPheHisThrTyrCysMe 258
167 AA.....ATCNCGTGATGACCAGTTTGACTTTTCATACATACTGTAT 128
258 tArgLysIleThrIleuArgSerTyrValAspLeuLeuLysLeuGluAspV 275
127 GAGGAGAGATTACCTTATAGATCATATGNGGACTTATTAAGAACTAGAAGATG 78
275 alLeuArgGlnHisProPheTyrPheLysAlaAlaArgIleAlaIleGlu 291
77 NACITCGACAGCATCCNTTTTACTTCAAGGCAGCAAGAAATTGCTATANAG 28

292 ileTyrLeuLysLeuHisAsp 298
27 ATCTATTGAAGCTTCATGAC 7

seq_name: /SIDS1/gcgdata/hold-geneseq/geneseq-emb1/NA1989.DAT.AAN90541

seq_documentation_block:
ID AAN90541 standard; DNA; 2703 BP.

XX AC AAN90541;
XX DT 28-NOV-1989 (first entry)
XX DE DNA encoding N-alpha-acetyl transferase.
XX KW N-alpha-acetyl transferase; herbicide resistance;
XX KW protein N-acetylation.
XX FH Key Location/Qualifiers
FT misc_feature 272..335 /*tag= a
FT misc_feature 338..392 /*tag= b
FT misc_feature 479..515 /*tag= c
FT misc_feature 542..566 /*tag= d
FT misc_feature 971..989 /*tag= e
FT misc_feature 1007..1049 /*tag= f
FT misc_feature 1061..1085 /*tag= g
FT misc_feature 1088..1130

seq_documentation_block:
ID AAH12222 standard; cDNA; 488 BP.
AC AAH12222;
XX DT 26-JUN-2001 (first entry)
XX DE Human cDNA clone (3'-primer) SEQ ID NO:9057.
XX KW Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.
XX OS Homo sapiens.
XX PN EP1074617-A2.
XX PD 07-FEB-2001.
XX PF 28-JUL-2000; 2000EP-0116126.
XX PR 29-JUL-1999; 99JP-0248036.
XX PR 27-AUG-1999; 99JP-0300253.
XX PR 11-JAN-2000; 2000JP-0118776.
XX PR 02-MAY-2000; 2000JP-0183767.
XX PR 09-JUN-2000; 2000JP-0241899.
XX PA (HELI-) HELIX RES INST.
XX PI Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
XX PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX DR WPI: 2001-318749/34.
XX PT Primer sets for synthesizing polynucleotides, particularly the 5602
XX PT full-length cDNAs defined in the specification, and for the detection
XX PT and/or diagnosis of the abnormality of the proteins encoded by the
XX PT full-length cDNAs -
XX PS Claim 3; SEQ ID 9057; 2537pp + CD ROM; English.

XX The present invention describes primer sets for synthesizing 5602
XX full-length cDNAs defined in the specification. Where a primer set
XX comprises: (a) an oligo-dr primer and an oligonucleotide complementary
XX to the complementary strand of a polynucleotide which comprises one of
XX the 5602 nucleotide sequences defined in the specification, where the
XX oligonucleotide comprises at least 15 nucleotides; or (b) a combination
XX of an oligonucleotide comprising a sequence complementary to the
XX complementary strand of a polynucleotide which comprises a 5'-end
XX sequence and an oligonucleotide comprising a sequence complementary to a
XX polynucleotide which comprises a 3'-end sequence, where the
XX oligonucleotide comprises at least 15 nucleotides and the combination of
XX the 5'-end sequence/3'-end sequence is selected from those defined in
XX the specification. The primer sets can be used in antisense therapy and
XX in gene therapy. The primers are useful for synthesizing polynucleotides,
XX particularly full-length cDNAs. The primers are also useful for the
XX detection and/or diagnosis of the abnormality of the proteins encoded by
XX cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
XX cDNAs 3633 to AAH18742 represent human cDNA sequences; AAB92446 to
XX AAH95893 represent human amino acid sequences; and AAH13629 to AAH13632
XX represent oligonucleotides, all of which are used in the exemplification
XX of the present invention.

SQ Sequence 488 BP; 121 A; 98 C; 84 G; 167 T; 18 other;

alignment_scores:
Quality: 605.50 Length: 157
Ratio: 4.388 Gaps: 1
Percent Similarity: 87.898 Percent Identity: 82.166
alignment_block:
US-09-836-410A-1 x AAH12222/rev ..

```

FT      misc_feature      /*tag= h
FT      1481..1505
FT      /*tag= i
FT      1829..1856
FT      misc_feature      /*tag= j
FT      1862..1885
FT      misc_feature      /*tag= k
FT      1909..1934
FT      /*tag= l
FT      misc_feature      2072..2117
FT      /*tag= m
FT      2123..2183
FT      /*tag= n
XX
XX      W08907138-A.
XX
XX      10-AUG-1989.
XX
XX      07-FEB-1989;      89WO-US00471.
XX
XX      08-FEB-1988;      88US-0153361.
XX      14-DEC-1988;      88US-0284344.
XX
XX      (GEO ) THE GENERAL HOSPITAL CORPORATION.
XX
XX      Smith JA, Lee FJS;
XX      WPI; 1989-249008/34.
XX      P-PSDB; AAP91070.
XX
XX      New pure N-alpha-acetyl transferase and DNA encoding it - catalysing
XX      acetylation of proteins and peptides, eg to stabilise pharmaceuticals
XX      or induce herbicide resistance in plants.
XX
XX      Claim 8: Page 50; fig 12b-e; 72pp; English.
XX
XX      DNA encodes N-alpha-actyl transferase, used for catalysing N-acetylation
XX      of peptides/proteins, eg to stabilise pharmaceuticals or to induce
XX      herbicide resistance in plants. Features a - n are fragments resulting
XX      from exonuclease III deletion. See also AAP91070.
XX
XX      Sequence 2703 BP; 943 A; 489 C; 530 G; 741 T; 0 other;

alignment_scores:
    Quality: 550.50      Length: 595
    Ratio: 1.605         Gaps: 17
    Percent Similarity: 57.647      Percent Identity: 27.059

alignment_block:
- US-09-836-410A-1 x AAN90541 ..
Align seg 1/1 to: AAN90541 from: 1 to: 2703

15  TyrProArgGlyLeuValProArgLysLeuProLeuAsnPheLeuSerG1 31
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
913  TATCCAAAGATGCGACCCACCACCAATTTATTCATTAACCTTCCCTCAAGA 962

31  Y...GluLysPheLysGluCysLeuAspArgPheLeuArgMetAsnPhes 47
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
963  CAAAGAAGAGCTCAGCAAAAATTCAGAGAATATGTTTGCCTCAATTGG 1012

47  erLysGlyCysProValPheAsnThrLeuArgSerLeuTyr...Arg 62
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
1013  AGCGGGTGTTCAGCAACTTTTCCAAAGTGAACCCCTTTACCAAGA 1062

63  AspLysGluLysValAla...IleValGluLysValValGlyTyrG1 78
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
1063  AGCAAGTCCAGGTTTCACCACTATTTGAGAAAATGTCCTTGATTATT 1112

78  uThrSerLeuLysSerCysArgLeuPheAsnProAsnAspGlyLysG 95
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
1113  GTCCGGATTA.....GATCCTACGAGGAT..... 1137

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95  luGluProProThrThrLeuLeuTyrValGlnTyrTyrLeuAlaGlnHis 111
||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
1138  .....CCAATTCCTTTTATTTGGACCAATTTACTTGCTCANCAT 1179

112  TyrAspLysIleGlyGlnProSerIleAlaLeuGluTyrIleAsnThrAl 128
::: ||| ||||| ||||| ||||| ||||| ||||| |||||
1180  TTCCTTTTCCTTAAGGATTTTCGAAAGCCCAAGAATATATTGATGCTGC 1229

128  aIleGluSerThrProThrLeuLeuPheLeuValLysAlaLysI 145
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
1230  CCTTGACCACACCCCACTTATGTTGAGTTTACATCCTCAAGGCAGTA 1279

145  leTyrLysHisAlaGlyAsnIleLysGluAlaAlaArgTyrMetAspGlu 161
|| ||||| ||| ||| ||||| ||||| ||||| ||||| |||||
1280  TCCTGAAGCACTTAGGCCTAATGGACACAGCGCTGGAATTTTGGAGGAA 1329

162  AlaGlnAlaLeuAspThrAlaAspArgPheIleAsnSerLysCysAlaLys 178
::: ||||| ||||| ||||| ||||| ||||| ||||| |||||
1330  GGTAGGCAACTTGATTTGCAGGATAGATTATCAACTGTAAACCGGTTAA 1379

178  sTyrMetLeuLysAlaAsnLeuIleLysGluAlaGluGluMetCysSerL 195
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
1380  GTACTTTTAAAGGCTAACAATATCGCAAGCGGTGGGANGTCGCGTCCC 1429

195  ysPheThrArg.....GluGlyThrSerAlaValGluAsnLeuAsnGlu 209
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
1430  TTTTCACCAAAACGATGATCTCTGTTAATGTTAAGGACTTACACCTT 1479

210  MetGlnCysMetTrpPheGlnThrGluCysAlaGlnAlaTyrLysAlaMe 226
::: ||||| ||||| ||||| ||||| ||||| ||||| |||||
1480  GTGGAGCTTCTTGTTTCATCGTAGAACAGGCGAAGCCTATTATAGACT 1529

226  t..... 226

1530  ATACCTGGATACAAAAGAAATTTAGACGATTTAGCATCGCTAAAAAAG 1579

226  ..... 226

1580  AGTTTCAAAGTGATAAAAGCGAACAAATTTGCGAATGATATCAAGAAAAAC 1629

227  .....AsnLysPhe...GlyGluAlaLeuLysLysCysHisG1 238
::: ||||| ||||| ||||| ||||| ||||| ||||| |||||
1630  CAATGGCTTGTCGCAATATTAAGTTTGGCGCTGAAAAGATTCAACGC 1679

238  ileGluArgHisPheIleGluIleThrAspAspGlnPheAspPheHisT 255
||| ::| ::| ::| ||||| ||||| ||||| ||||| |||||
1680  TATTCCAAAGTTTATAAACAATTCGAAGATGACCAGTTGGATTTCATT 1729

255  hrTyrCysMetArgLysIleThrLeuArgSerTyrValAspLeuLys 271
::: ||||| ||||| ||||| ||||| ||||| ||||| |||||
1730  CATACTGTATGAGAAAAGGTACGCCAAGAGCCTATCTGGAGATGTTAGA 1779

272  LeuGluAspValLeuArgGlnHisProPheTyrPheLysAlaAlaArgI1 288
||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
1780  TGGGAAAAGGCACITTTATACCAACCCCATGATGTTCCGCGCAATGAAGGA 1829

288  eAlaIleGluIleTyrLeuLysLeuHisAspAsnProLeuThrAspGluA 305
||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
1830  AGCATCAAAGCTTTACTTTCAATGCAATGATGATCGCTTA..... 1869

305  snLysGluHisGluAlaAspThrAlaAsnMetSerAspLysGluLys 321
::: ||||| ||||| ||||| ||||| ||||| ||||| |||||
1870  .....AAAAGAAAGTCCGATTTCTTTAGATGAAAATTCAGATGAAATCAA 1914

322  LysLeuArgAsnLysGlnArgAlaGlnLysLysAlaGlnIleGluG1 338
::: ||||| ||||| ||||| ||||| ||||| ||||| |||||
1915  AATAATGCCAAAATAGTAGCAGCCAAAAGAAAACCTAAGAAGGAGGC 1964

338  uGluLysLysAsnAlaGluLysGluLysProGlnArgAsnProLysLysL 355
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Mon Jul 22 09:40:54 2002

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 DEFINITION Sequence 1 from Patent WO01/79506.
 ACCESSION AX285242
 VERSION AX285242.1 GI:17045930

KEYWORDS
 SOURCE human.

ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE
 1 (sites)
 Gendron, R.L. and Paradis, H.
 Treatment of ocular neovascularization and related diseases
 Patent: WO 0179506-A 1 25-OCT-2001;
 Children's Hospital Research Foundation (US)

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VERSION     AX285294.1  GI:17045975
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            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1 (sites)
AUTHORS     Gendron,R.L. and Paradis,H.
TITLE       Inhibition of bone tumor formation using antisense cdna therapy
JOURNAL     Patent: WO 0179505-A 2 25-OCT-2001;
            CHILDREN'S HOSPITAL MEDICAL CENTER (US)
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DEFINITION Sequence 4 from Patent WO01/9505.

ACCESSION AX285296

VERSION AX285296.1 GI:17045977

KEYWORDS human.

SOURCE

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (sites)

Gendron, R.L. and Paradis, H.

TITLE Inhibition of bone tumor formation using antisense cdna therapy

JOURNAL Patent: WO 01/9505-A 4 25-OCT-2001;

CHILDREN'S HOSPITAL MEDICAL CENTER (US)

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Location/Qualifiers

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Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
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Gendron,R.L., Adams,L.C. and Paradis,H.
Tubedown-1, A novel acetyltransferase associated with blood vessel
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Dev. Dyn. 118 (2), 300-315 (2000)
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2 (bases 1 to 3421)
Gendron,R.L., Adams,L.C. and Paradis,H.
Direct Submission
Submitted (20-FEB-2000) Pediatrics, Childrens Hospital Medical
Center, 3333 Burnet Avenue, Cincinnati, OH 45229-3039, USA
3 (bases 1 to 3421)
Gendron,R.L., Adams,L.C. and Paradis,H.
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Submitted (13-JUN-2000) Pediatrics, Childrens Hospital Medical
Center, 3333 Burnet Avenue, Cincinnati, OH 45229-3039, USA
Amino acid sequence updated by submitter
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ACCESSION AF237622
VERSION AF237622.1 GI:8164012
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DEFINITION Homo sapiens putative acetyltransferase mRNA, complete cds.

ACCESSION AF327722

VERSION AF327722.1 GI:13195459

KEYWORDS

SOURCE human.

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Cranial; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 4192)

AUTHORS He, Y. G., Tan, D. Y., Lai, J. H., Xie, Y. F. and Qian, W.

TITLE Cloning and analysis of a human putative acetyltransferase

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 4192)

AUTHORS He, Y. G., Tan, D. Y., Lai, J. H., Xie, Y. F. and Qian, W.

TITLE Direct Submission

JOURNAL Submitted (11-DEC-2000) Biology Department, Yunnan University,
North Street of Greenlake, Kunming, Yunnan 650091, China
FEATURES Location/Qualifiers
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VERSION AJ314788.1 GI:14589341
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SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
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Fluge,O., Bruland,O., Akslen,L.A., Varhaug,J.E. and Lillehaug,J.R.
Identification of NATH, a novel gene overexpressed in papillary
thyroid carcinomas
Unpublished
2 (bases 1 to 5505)
Fluge,O.
Direct Submission
Submitted (31-MAY-2001) Fluge O., Dept. of Molecular Biology,
University of Bergen, Thormohlens gt 55, N-5020 Bergen, NORWAY
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VERSION AF247679.1 GI:9651962
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Amphibia; Batrachia; Anura; Mesobatrachia; Pipoidae; Pipidae;
Xenopodinae; Xenopus.
REFERENCE
1 (bases 1 to 3324)
Choi,S.-C., Kim,J. and Han,J.-K.
Expression of N-terminal acetyltransferase in Xenopus laevis
Unpublished
JOURNAL
REFERENCE
2 (bases 1 to 3324)
Choi,S.-C., Kim,J. and Han,J.-K.
Direct Submission
JOURNAL
Submitted (22-MAR-2000) Life Science, Pohang University of Science
and Technology, Pohang 790-784, South Korea
FEATURES
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oligo capping; fis (full insert sequence).
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 Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.

REFERENCE 1 (sites)
 AUTHORS Isogai,T., Ota,T., Hayashi,K., Sugiyama,T., Otsuki,T., Suzuki,Y.,
 Nishikawa,T., Nagai,K., Sugano,S., Shiratori,A., Sudo,H.,
 Wagatsuma,M., Hosoiri,T., Kaku,Y., Kodaira,H., Kondo,H.,
 Sugawara,M., Takahashi,M., Chiba,Y., Ishida,S., Murakawa,K.,
 Ono,Y., Takiguchi,S., Watanabe,S., Kimura,K., Murakami,K.,
 Ishii,S., Kawai,Y., Saito,K., Yamamoto,J., Wakamatsu,A.,
 Nakamura,Y., Nagahara,K., Masuho,Y., Ninomiya,K. and Iwayanagi,T.
 NEDO human cDNA sequencing project
 Unpublished (2000)

JOURNAL 2 (bases 1 to 1802)
 REFERENCE Isogai,T. and Otsuki,T.
 AUTHORS Direct Submission
 TITLE Submitted
 JOURNAL Submitted (23-AUG-2000) to the DDBJ/EMBL/GenBank databases. Takao
 Isogai, Helix Research Institute, Genomics Laboratory; 1532-3 Yana,
 Kisarazu, Chiba 292-0812, Japan (E-mail: genomics@hri.co.jp,
 Tel:81-438-52-3951, Fax:81-438-52-3952)

COMMENT NEDO human cDNA sequencing project supported by Ministry of
 International Trade and Industry of Japan; cDNA full insert
 sequencing; Research Association for Biotechnology; cDNA library
 construction, 5'- & 3'-end one pass sequencing and clone selection;;
 Helix Research Institute (supported by Japan Key Technology Center
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 University of Tokyo.

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 1 (sites)
 Isogai, T., Ota, T., Hayashi, K., Sugiyama, T., Otsuki, T., Suzuki, Y.,
 Nishikawa, T., Nagai, K., Sugano, S., Shiratori, A., Sudo, H.,
 Wagatsuma, M., Hosohashi, T., Kaku, Y., Kodaira, H., Kondo, H.,
 Ono, Y., Takiguchi, S., Chiba, Y., Ishida, S., Murakami, K.,
 Ota, Y., Takiguchi, S., Watanabe, S., Kimura, K., Murakami, K.,
 Ishii, S., Kawai, Y., Saito, K., Yamamoto, J., Wakamatsu, A.,
 Nakamura, Y., Nagahara, K., Masuho, Y., Ninomiya, K. and Iwayanagi, T.
 NEDO human cDNA sequencing project
 Unpublished (2000)
 2 (bases 1 to 1985)
 Isogai, T. and Otsuki, T.
 Direct Submission
 Submitted (23-AUG-2000) Takao Isogai, Helix Research Institute,
 Genomics Laboratory, 1532-3 Yana, Kisarazu, Chiba 252-0812, Japan
 (E-mail: genomics@hri.co.jp, Tel: 81-438-52-3951, Fax: 81-438-52-3952)
 NEDO human cDNA sequencing project supported by Ministry of
 International Trade and Industry of Japan; cDNA full insert
 sequencing: Research Association for Biotechnology; cDNA library
 construction, 5'- & 3'-end one pass sequencing and clone selection;
 Helix Research Institute (supported by Japan Key Technology Center
 etc.) and Department of Virology, Institute of Medical Science,
 University of Tokyo.
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1 (bases 1 to 180213)
Celiner,S.E., Adams,M.D., Kronmiller,B., Tyler,D., Wan,K.H.,
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Rogers,Y., An,H., Baldwin,D., Banazon,J., Beeson,K.Y., Busam,D.A.,
Carlson,J.W., Center,A., Champe,M., Davenport,L.B., Dietz,S.M.,
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Feria,S., Frise,E., Galle,R.F., Garg,N.S., George,R.A.,
Gonzalez,M., Houck,J., Hoskins,R.A., Hostin,D., Howland,T.J.,
Ibegwam,C., Jalali,M., Kruse,D., Li,P., Mattel,B., Moshrefi,A.,
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Pacleb,J., Paragas,V., Park,S., Patel,S., Pfeiffer,B.,

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TITLE
JOURNAL
REFERENCE
AUTHORS
Phouanavong,S., Pittman,G.S., Puri,V., Richards,S., Scheeler,F.,
Stapleton,M., Strong,R.O., Svirskas,R., Tector,C., Williams,S.M.,
Zaveri,J.S., Smith,H.O., Rubin,G.M. and Venter,J.C.
Sequencing of Drosophila chromosome X, region 18D-18D
Unpublished
2 (bases 1 to 180213)
Celiner,S.E., Agbayani,A., Arcaina,T.T., Baxter,E., Blazej,R.G.,
Butenhoff,C., Champe,M., Chavez,C., Chew,M., Ciesiolka,L.,
Doyle,C.M., Farfan,D.E., Galle,R., George,R.A., Harris,N.L.,
Hoskins,R.A., Houston,K.A., Hummasti,S.R., Karra,K., Kearney,L.,
Kim,E., Lee,B., Lewis,S., Li,P., Lomont,M.A., Mazda,P.,
Moshrefi,A.R., Moshrefi,M., Nixon,K., Pacleb,J.M., Park,S.,
Pfeiffer,B., Poon,L., Sequeira,A., Sethi,H., Snir,E.,
Svirskas,R.R., Wan,K.H., Weinburg,T., Zhang,R., Zieran,L.L. and
Rubin,G.M.
Direct Submission
Submitted (24-SEP-1999) Drosophila Genome Center, Lawrence Berkeley
Laboratory, MS 64-121, Berkeley, CA 94720, USA
On Mar 17, 2001 this sequence version replaced gi:6563418.
Sequence submitted by:
Berkeley Drosophila Genome Project
Lawrence Berkeley National Laboratory, MS 64-121
Berkeley, CA 94720
This sequence was assembled using end sequences from a whole genome
shotgun and from subclones of this BAC and its neighboring clones.
For further information about this sequence, including its location
and relationship to other sequences, please visit our sequence
archive Web site (http://www.fruitfly.org/sequence/) or send email
to bdgp@fruitfly.berkeley.edu.
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:::|||||:::|||||:::|||||:::|||||:::|||||
24830 GCAGGAGGCCCAAAATCTACACAGCTTGGCGGACGCGCATGTGTTTG 24879
544CysLysGluAlaAlaGluAlaTyrArgAlaSerCysHisLysLeu 558
:::|||||:::|||||:::|||||:::|||||:::|||||
24880 GCGAGTGGCGAGGAGCGCGCTCTCTACCAGCAGCGCTGCCCAACCGC 24929
559 PheProTyrAlaLeuAlaPhe..... 565
|||||:::|||||:::|||||:::|||||:::|||||
24930 TTCCAGTACGCCCGCATCTTTCCGCAATCTCGAGGAGCTGGAGGCTCACT 24979
566MetProGlyTyrGluGluAspMetLysIleT 577
|||||:::|||||:::|||||:::|||||:::|||||
24980 GCAGGAGAGGAGGCGCCAAAGCTGCGCGCGGAGAGAGCAGCAGCAGCAGC 25029
577 hrValAsnGlyAspSerSerAlaGluThrGluGluLeuAla 590
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25030 TGATTACGTTGATCTAGTGAACCGGTGTCTGTTTGGCC 25070

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seq_documentation_block:
LOCUS      AC011071                      182623 bp    DNA        linear        INV 30-MAY-2001
DEFINITION Drosophila melanogaster, chromosome X, region 18C-18D, BAC clone
BACR27L16, complete sequence.
ACCESSION  AC011071
VERSION    AC011071.12  GI:14249062
SOURCE     fruit fly.
ORGANISM   Drosophila melanogaster
            Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
            pterygota; Endopterygota; Diptera; Brachycera;
            Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 182623)
Celisner, S.E., Adams, M.D., Kronmiller, B., Tyler, D., Wan, K.H.,
Holt, R.A., Evans, C.A., Gocayne, J.D., Amanatides, P.G., Brandon, R.C.,
Rogers, Y., An, H., Baldwin, D., Banzon, J., Beeson, K.Y., Busam, D.A.,
Carlson, J.W., Center, A., Champs, M., Davenport, L.B., Dietz, S.M.,
Dodson, K., Dorsett, V., Dou, L.E., Doyle, C., Dresnek, D., Farfan, D.,
Ferriera, S., Frise, E., Galle, R.F., Garg, N.S., George, R.A.,
Gonzalez, M., Houck, J., Hoskins, R.A., Hostin, D., Howland, T.J.,
Ibegwam, C., Jalali, M., Kruse, D., Li, P., Mattei, B., Moshrefi, A.,
McIntosh, T.C., Moy, M., Murphy, B., Nelson, C., Nelson, K.A., Nunoo, J.,
Pacleb, J., Paragas, V., Park, S., Patel, S., Pfeiffer, B., Scheeler, F.,
Phouanavong, S., Pittman, G.S., Puri, V., Richards, S., Scheeler, F.,
Stapleton, M., Strong, R., Svirska, R., Tector, C., Williams, S.M.,
Zaveri, J.S., Smith, H.O., Rubin, G.M. and Venter, J.C.
Sequencing of Drosophila chromosome X, region 18C-18D
Unpublished
2 (bases 1 to 182623)
Celisner, S.E., Agbanyani, A., Arcaluna, T.T., Baxter, E., Blazej, R.G.,
Butenhorff, C., Champs, M., Chavez, C., Chew, M., Ciesiolka, L.,
Doyle, C.M., Farfan, D.E., Galle, R., George, R.A., Harris, N.L.,
Hoskins, R.A., Houston, K.A., Hummasti, S.R., Karra, K., Kearney, L.,
Kim, E., Lee, B., Lewis, S., Li, P., Lomotan, M.A., Mazda, P.,
Moshrefi, A.R., Moshrefi, M., Nixon, K., Pacleb, J.M., Park, S.,
Pfeiffer, B., Poon, L., Sequeira, A., Sethi, H., Snir, E.,
Svirska, R.R., Wan, K.H., Weinburg, T., Zhang, R., Zieran, L.L. and
Rubin, G.M.
Direct Submission
Submitted (01-OCT-1999) Drosophila Genome Center, Lawrence Berkeley
Laboratory, MS 64-121, Berkeley, CA 94720, USA
On May 30, 2001 this sequence version replaced gi:7143400.
Sequence submitted by:
Berkeley Drosophila Genome Project
Lawrence Berkeley National Laboratory, MS 64-121
Berkeley, CA 94720
This sequence was assembled using end sequences from a whole genome
shotgun and from subclones of this BAC and its neighboring clones.
For further information about this sequence, including its location
and relationship to other sequences, please visit our sequence
archive Web site (http://www.fruitfly.org/sequence/) or send email
to bdgp@fruitfly.berkeley.edu.
Location/Qualifiers
1..182623
/organism="Drosophila melanogaster"
/strain="y; cn bw sp"
/db_xref="taxon:7227"
/chromosome="x"
/map="18C-18D"
/clone="BACR27L16 (D903)"
/clone_lib="RPC1-98 (Roswell Park Cancer Institute
Drosophila melanogaster BAC library, partial EcoRI in
PBACe3.6)"
BASE COUNT  49691 a 41087 c 41478 g 50367 t
ORIGIN
alignment_scores:
Quality: 1385.00
Ratio: 2.934
Percent Similarity: 69.310
Length: 681
Gaps: 12
Percent Identity: 42.291

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```

alignment_block:
US-09-836-410A-1 x AC011071
Align seg 1/1 to: AC011071 from: 1 to: 182623
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: : : : : : : : : : : : : : : : : : : : : : : : : :
84550 GTGTCGATCTAICGGCTGTTCAGGAGCAGTATCCGCGCGCTGTGCC 84599
: : : : : : : : : : : : : : : : : : : : : : : : : :
21 OArgLysLeuProLeuAsnPhenLeuSerGlyGluLysPheLysGluCysL 38
: : : : : : : : : : : : : : : : : : : : : : : : : :
84600 TCGCGCGCTGCTTTGACATCGCAACGCGCAGGTTTCGCGCTGCGA 84649
: : : : : : : : : : : : : : : : : : : : : : : : : :
38 eAspArgPheLeuArgMetAsnPhenSerLysGlyCysProProValPhe 54
: : : : : : : : : : : : : : : : : : : : : : : : : :
84650 CCGACGAGTACCTTCGCGCGCTGCTCGTAAGGGCATTTCGCGCTATT 84699
: : : : : : : : : : : : : : : : : : : : : : : : : :
55 AsnThrLeuArgSerLeuTyrArgAspLysGluLysValAlaIleValG 71
: : : : : : : : : : : : : : : : : : : : : : : : : :
84700 GTCAACGTGGCGACTCTGCACACAGATACCGGAGGCGCGCTTATCGA 84749
: : : : : : : : : : : : : : : : : : : : : : : : : :
71 uGluLeuValValGlyTyrGluThrSerLeuLysSerCysArgLeuPhe 88
: : : : : : : : : : : : : : : : : : : : : : : : : :
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: : : : : : : : : : : : : : : : : : : : : : : : : :
88 sProAsnAspAspGly.....LysGluGluProProThrThrLeu 101
: : : : : : : : : : : : : : : : : : : : : : : : : :
84800 CTCGGAAGATGCGGCGCGCGGATTCCTCCGTCGAGCGCGCTCGGCGTG 84849
: : : : : : : : : : : : : : : : : : : : : : : : : :
102 LeuTrpValGlnTyrTyrLeuAlaGlnHisTyrAspLysIleGlyGlnPr 118
: : : : : : : : : : : : : : : : : : : : : : : : : :
84850 GTGTGGACGCGCTGTTCTTGGCGCAGCAGTACGACTACATCGCGCATC 84899
: : : : : : : : : : : : : : : : : : : : : : : : : :
118 oSerIleAlaLeuGluTyrIleAsnThrAlaIleGluSerThrProThrL 135
: : : : : : : : : : : : : : : : : : : : : : : : : :
84900 GGACGCGCTCTGGAGTACATCAATGTGGTATCGACCATACGCCACAC 84949
: : : : : : : : : : : : : : : : : : : : : : : : : :
135 eGluLeuLeuPheLeuValLysAlaLysIleTyrLysHisAlaGlyAsn 151
: : : : : : : : : : : : : : : : : : : : : : : : : :
84950 TCATTGAGCTGCTTATCACCAGGTCGCATCTTAAAGCATGCTGGCGAT 84999
: : : : : : : : : : : : : : : : : : : : : : : : : :
152 IleLysGluAlaAlaArgTrpMetAspGluAlaGlnAlaLeuAspThrAl 168
: : : : : : : : : : : : : : : : : : : : : : : : : :
85000 CCCGTGGAGGCGTACGCTGTGCTGGAGGAGGCCCAAGCATGACACGCG 85049
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: : : : : : : : : : : : : : : : : : : : : : : : : :
85050 AGATCGGTGAGTGAGCCCGCTGAGTACATAGACCTGCTGAACCATGTA 85099
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: : : : : : : : : : : : : : : : : : : : : : : : : :
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: : : : : : : : : : : : : : : : : : : : : : : : : :
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: : : : : : : : : : : : : : : : : : : : : : : : : :
85150 ATGCTGCGCGCCAACTGGTCGAGGAGGCGAGGAGATCTCGGCCAAGTT 85199
: : : : : : : : : : : : : : : : : : : : : : : : : :
196 eThrArgGluGlyThrSerAlaValGluAsnLeuAsnGluMetGlnCysM 213
: : : : : : : : : : : : : : : : : : : : : : : : : :
85200 CACTCGCGAGGCTGTCTCCGCCATGGACAACCTTGAACGAGATGCGATGA 85249
: : : : : : : : : : : : : : : : : : : : : : : : : :
213 eTrpPheGlnThrGluCysAlaGlnAlaTyrLysAlaMetAsnLysPhe 229
: : : : : : : : : : : : : : : : : : : : : : : : : :
85250 TGTGGTTCCAGACGGAGTGTGCCCTTGGCCATGACGCGCATGGTGGCTGG 85299
: : : : : : : : : : : : : : : : : : : : : : : : : :
230 GlyGluAlaLeuLysCysHisGluIleGluArgHisPheIleGluL 246
: : : : : : : : : : : : : : : : : : : : : : : : : :
85300 GCGGAGTCTGTGAAGAGTGCACGAGGTGGAGCGCCACTTTTGGCCGAAT 85349
: : : : : : : : : : : : : : : : : : : : : : : : : :
246 eThrAspAspGlnPheAspPheHisThrTyrCysMetArgLysIleThrL 263
: : : : : : : : : : : : : : : : : : : : : : : : : :
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: : : : : : : : : : : : : : : : : : : : : : : : : :

```


GKSTQPOXYLVDFGYSBIACTOPBRLACYSICKRVVAHELDDYSGSRVAFQIRPERSKT
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 KLIMSTINVELFGFEGEARLVGVRPLRFLKRYLPAPPELLAKQAQTSKQK
 PLNIDPAFVOVLSDQQYPTSEKRGDVLFGVSNELSVSEVHEVYATQTHMLFAT
 RHSGQADQKVDYAPBEGMKCTVSTNAETSLVDGVFVDSGVKYEKNWFAT
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alignment_scores:
  Quality: 1385.00      Length: 681
  Ratio: 2.934          Gaps: 12
  Percent Similarity: 69.310      Percent Identity: 42.291

alignment_block:
  US-09-836-410A-1 x AE003512      ..

Align seg 1/1 to: AE003512 from: 1 to: 301457

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[illegible]

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230 GlyGluAlaLeuLysLysCysHisGluIleGluAtrGHisPheIleGluI 246
GGCGAGTCGTTGAAGAGTCCACGAGGTGGAGGCCCACTTTGCCGAAT 843559

246 éthrAspAspGlnPheAspPheHisThrTyrCysMetArgLysIleThrL 263
CGTCGAGGACCAAGTTCGATTCCACACCATATTGCATCGTAAATGACGC 844508

263 euArgSerTyrValAspLeuLeuLysLeuGluAspValLeuArgGlnHis 279
TCGCGCCCTACGCTTGGCTGTGGATTGGAGGAGCTGCTACGCCAGCAT 844559

280 ProPheTyrPheLysAlaAlaArgIleAlaIleGluIleTyrLeuLysLe 296
CCATTCTATTCAAGCGGCCAAGTCGCCCATCGAGGTGTACATTCGTCT 84509

296 uHisAspAsnProLeuThrAspGluAsnLysGluHisGluAlaAsp... 311
GTCACGACAGCGCTTAAGTCGGAGACACCACTTGAAGAGATTCACATTCG 84559

311 :..... 311

84609 GTATGTGGACATTACCCCACTAGCTCCCTGCTGCTTCTAGTTAGCTGTA 84658

312 :.....ThrAlaAsnMetSerAspLysGluLeuLysLysLy 322
TGACCGCCCTTCTTCTGTTACAGAGAACCCTGCCGCATCAGAACTGAAGA 846708

322 sLeuArgAsnLysGlnArgAlaGlnLysLysAlaGlnIleGluGlu. 338
GCTGCGCAGCAACAGCCGACAGGCCAAGAAAGCTGAGCTGGAGAGTG 84708

339 :GluLysLysAsnAlaGluLysGluLysProGlnArgAsnProLysLys 354
CGCAGCGCGCACAGCGCAGCTGAAGCGCGAGCAGCACCAGAAATCGAA 84808

355 LysLysAspAspAspGluGluIleGlyClyProLys...GluGluLe 370
CAGCAGCAACACGAGGAGCAGCCCGCATGCTCCGAGTTGGACGAGCT 84808

370 uIleProGluLysLeuAlaLysValGluThrProLeuGluGluAlaIleL 387
GGTGGCCCGAGAAGCTGGAGCGCACGAGCATCCCTGGCAGAGCCCATTCG 84859

387 ysPheLeuThrProLeuLysAsnLeuValLysAsnLysIleGluThrHis 403
AATTCTTGAACCCGCTCGACGAGCTGGCTAAGGAGCGCATCGAGAGCAT 84908

404 LeuPheAlaPheGluIleTyrPheArgLysGluLysPheLeuLeuMetLe 420
CTGCTGGCTTCGAGTGTACTACCGCAGAACAACTGCTCCTTATGCT 84958

420 uGlnSerValLysArgAlaPheAlaIleAspSerSerHisProTrpLeuH 437
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8509 CGAGTGCATTCGGCTTGGTTAAATCGTTTACCAGCGCCGCAAGGAG 85058

437 IsGluCysMetIleArgLeuPheHisSerValCysGlu...SerLysAsp 452
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453 LeuPro.....GluThrValArgThrValLeuLysGlnGluMetAsnAr 467
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467 gLeuPheGlyAlaThrAsnProLysAsnPheAsnGluThrPheLeuLysA 484
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Mon Jul 22 09:40:54 2002

```
484 rgAsnSerAspSerLeuProHisArgLeuSerAlaAlaLysMetValTyr 500
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85209 AACACAAGCTTCCATATGCTATACGAGGCGCACGAGTCTGTAC 85258
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517 uAsp.....GlySerLeuT 522
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522 hrAsnArgAsnLeuGlnThrCysMetGluValLeuGluAlaLeuCysAsp 538
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539 Gly.....Ser.LeuArgAsp..... 543
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559 PheProTyrAlaLeuAlaPhe..... 565
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